APPLICATION FOR PATENT

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Title:

POLYNUCLEOTIDE ENCODING A POLYPEPTIDE HAVING HEPARANASE ACTIVITY AND EXPRESSION OF SAME IN GENETICALLY MODIFIED CELLS

This is a continuation of U.S. Patent Application No. 09/776,874, filed February 6, 2001, which is a continuation of U.S. Patent Application No. 09/258,892, filed March 1, 1999, which is a continuation-in-part of PCT/US98/17954, filed August 31, 1998, which claims priority from U.S. Patent Application 09/109,386, filed July 2, 1998, now abandoned, which is a continuation-in-part of U.S. Patent Application 08/922,170, filed September 2, 1997, now, U.S. Patent No. 5,968,822.

FIELD AND BACKGROUND OF THE INVENTION

The present invention relates to a polynucleotide, referred to hereinbelow as *hpa*, encoding a polypeptide having heparanase activity, vectors (nucleic acid constructs) including same and genetically modified cells expressing heparanase. The invention further relates to a recombinant protein having heparanase activity and to antisense oligonucleotides, constructs and ribozymes for down regulating heparanase activity. In

Heparan sulfate proteoglycans: Heparan sulfate proteoglycans (HSPG) are ubiquitous macromolecules associated with the cell surface and extra cellular matrix (ECM) of a wide range of cells of vertebrate and invertebrate tissues (1-4). The basic HSPG structure includes a protein core to which several linear heparan sulfate chains are covalently attached. These polysaccharide chains are typically composed of repeating hexuronic and D-glucosamine disaccharide units that are substituted to a varying extent with N- and O-linked sulfate moieties and N-linked acetyl groups (1-4). Studies on the involvement of ECM molecules in cell attachment, growth and differentiation revealed a central role of HSPG in embryonic morphogenesis, angiogenesis, neurite outgrowth and tissue repair (1-5). HSPG are prominent components of blood vessels (3). In large blood vessels they are concentrated mostly in the intima and inner media, whereas in capillaries they are found mainly in the subendothelial basement membrane where they support proliferating and migrating endothelial cells and stabilize the structure of the capillary wall. The ability of HSPG to interact with ECM macromolecules such as collagen, laminin and fibronectin, and with different attachment sites on plasma membranes suggests a key role for this proteoglycan in the self-assembly and insolubility of ECM components, as well as in cell adhesion and

result in degradation of the subendothelial ECM and hence may play a

decisive role in extravasation of blood-borne cells. HS catabolism is observed in inflammation, wound repair, diabetes, and cancer metastasis, suggesting that enzymes which degrade HS play important roles in pathologic processes. Heparanase activity has been described in activated immune system cells and highly metastatic cancer cells (6-8), but research has been handicapped by the lack of biologic tools to explore potential causative roles of heparanase in disease conditions.

Involvement of Heparanase in Tumor Cell Invasion and Metastasis: Circulating tumor cells arrested in the capillary beds of different organs must invade the endothelial cell lining and degrade its underlying basement membrane (BM) in order to invade into the extravascular tissue(s) where they establish metastasis (9, 10). Metastatic tumor cells often attach at or near the intercellular junctions between adjacent endothelial cells. Such attachment of the metastatic cells is followed by rupture of the junctions, retraction of the endothelial cell borders and migration through the breach in the endothelium toward the exposed underlying BM (9). Once located between endothelial cells and the BM, the invading cells must degrade the subendothelial glycoproteins and proteoglycans of the BM in order to migrate out of the vascular compartment. Several cellular enzymes (e.g., collagenase IV, plasminogen

-D-glucuronidase (heparanase) that cleaves HS at specific intrachain sites (6, 8, 11). Expression of a HS degrading heparanase was found to correlate with the metastatic potential of mouse lymphoma (11), fibrosarcoma and melanoma (8) cells. Moreover, elevated levels of heparanase were detected in sera from metastatic tumor bearing animals and melanoma patients (8) and in tumor biopsies of cancer patients (12).

The control of cell proliferation and tumor progression by the local microenvironment, focusing on the interaction of cells with the extracellular matrix (ECM) produced by cultured corneal and vascular endothelial cells, was investigated previously by the present inventors. This cultured ECM closely resembles the subendothelium *in vivo* in its morphological appearance and molecular composition. It contains collagens (mostly type III and IV, with smaller amounts of types I and V), proteoglycans (mostly heparan sulfate- and dermatan sulfate- proteoglycans, with smaller amounts of chondroitin sulfate proteoglycans), laminin, fibronectin, entactin and elastin (13, 14). The ability of cells to degrade HS in the cultured ECM was studied by allowing cells to interact with a metabolically sulfate labeled ECM, followed by gel filtration (Sepharose 6B) analysis of degradation products released into the culture medium (11). While intact HSPG are eluted next to the void volume of the column

are eluted more toward the V_t of the column (0.5< kav< 0.8, Mr ~5-7x10^3) (11).

The heparanase inhibitory effect of various non-anticoagulant species of heparin that might be of potential use in preventing extravasation of blood-borne cells was also investigated by the present inventors. Inhibition of heparanase was best achieved by heparin species containing 16 sugar units or more and having sulfate groups at both the N and O positions. While O-desulfation abolished the heparanase inhibiting effect of heparin, O-sulfated, N-acetylated heparin retained a high inhibitory activity, provided that the N-substituted molecules had a molecular size of about 4,000 daltons or more (7). Treatment of experimental animals with heparanase inhibitors (e.g., non-anticoagulant species of heparin) markedly reduced (~90%) the incidence of lung metastases induced by B16 melanoma, Lewis lung carcinoma and mammary adenocarcinoma cells (7, 8, 16). Heparin fractions with high and low affinity to anti-thrombin III exhibited a comparable high anti-metastatic activity, indicating that the heparanase inhibiting activity of heparin, rather than its anticoagulant activity, plays a role in the anti-metastatic properties of the polysaccharide (7).

Heparanase activity in the urine of cancer patients: In an attempt

were screened (16a). Heparanase activity was detected in the urine of some, but not all, cancer patients. High levels of heparanase activity were determined in the urine of patients with an aggressive metastatic disease and there was no detectable activity in the urine of healthy donors.

Heparanase activity was also found in the urine of 20% of normal and microalbuminuric insulin dependent diabetes mellitus (IDDM) patients, most likely due to diabetic nephropathy, the most important single disorder leading to renal failure in adults.

Possible involvement of heparanase in tumor angiogenesis: Fibroblast growth factors are a family of structurally related polypeptides characterized by high affinity to heparin (17). They are highly mitogenic for vascular endothelial cells and are among the most potent inducers of neovascularization (17, 18). Basic fibroblast growth factor (bFGF) has been extracted from the subendothelial ECM produced *in vitro* (19) and from basement membranes of the cornea (20), suggesting that ECM may serve as a reservoir for bFGF. Immunohistochemical staining revealed the localization of bFGF in basement membranes of diverse tissues and blood vessels (21). Despite the ubiquitous presence of bFGF in normal tissues, endothelial cell proliferation in these tissues is usually very low, suggesting that bFGF is somehow sequestered from its site of action. Studies on the

20, 22). It was demonstrated that heparanase activity expressed by platelets, mast cells, neutrophils, and lymphoma cells is involved in release of active bFGF from ECM and basement membranes (23), suggesting that heparanase activity may not only function in cell migration and invasion, but may also elicit an indirect neovascular response. These results suggest that the ECM HSPG provides a natural storage depot for bFGF and possibly other heparin-binding growth promoting factors (24, 25). Displacement of bFGF from its storage within basement membranes and ECM may therefore provide a novel mechanism for induction of neovascularization in normal and pathological situations.

Recent studies indicate that heparin and HS are involved in binding of bFGF to high affinity cell surface receptors and in bFGF cell signaling (26, 27). Moreover, the size of HS required for optimal effect was similar to that of HS fragments released by heparanase (28). Similar results were obtained with vascular endothelial cells growth factor (VEGF) (29), suggesting the operation of a dual receptor mechanism involving HS in cell interaction with heparin-binding growth factors. It is therefore proposed that restriction of endothelial cell growth factors in ECM prevents their systemic action on the vascular endothelium, thus maintaining a very low rate of endothelial cells turnover and vessel growth. On the other hand,

processes such as wound healing, inflammation and tumor development (24, 25).

Expression of heparanase by cells of the immune system: Heparanase activity correlates with the ability of activated cells of the immune system to leave the circulation and elicit both inflammatory and autoimmune responses. Interaction of platelets, granulocytes, T and B lymphocytes, macrophages and mast cells with the subendothelial ECM is associated with degradation of HS by a specific heparanase activity (6). The enzyme is released from intracellular compartments (e.g., lysosomes, specific granules, etc.) in response to various activation signals (e.g., thrombin, calcium ionophore, immune complexes, antigens, mitogens, etc.), suggesting its regulated involvement in inflammation and cellular immunity.

Some of the observations regarding the heparanase enzyme were reviewed in reference No. 6 and are listed hereinbelow:

First, a proteolytic activity (plasminogen activator) and heparanase participate synergistically in sequential degradation of the ECM HSPG by inflammatory leukocytes and malignant cells.

Second, a large proportion of the platelet heparanase exists in a latent form, probably as a complex with chondroitin sulfate. The latent enzyme is activated by tumor cell-derived factor(s) and may then facilitate cell

Third, release of the platelet heparanase from α-granules is induced by a strong stimulant (i.e., thrombin), but not in response to platelet activation on ECM.

Fourth, the neutrophil heparanase is preferentially and readily released in response to a threshold activation and upon incubation of the cells on ECM.

Fifth, contact of neutrophils with ECM inhibited release of noxious enzymes (proteases, lysozyme) and oxygen radicals, but not of enzymes (heparanase, gelatinase) which may enable diapedesis. This protective role of the subendothelial ECM was observed when the cells were stimulated with soluble factors but not with phagocytosable stimulants.

Sixth, intracellular heparanase is secreted within minutes after exposure of T cell lines to specific antigens.

Seventh, mitogens (Con A, LPS) induce synthesis and secretion of heparanase by normal T and B lymphocytes maintained *in vitro*. T lymphocyte heparanase is also induced by immunization with antigen *in vivo*.

Eighth, heparanase activity is expressed by pre-B lymphomas and B-lymphomas, but not by plasmacytomas and resting normal B lymphocytes.

enzyme into the incubation medium. Similar results were obtained with human myeloid leukemia cells induced to differentiate to mature macrophages.

Tenth, T-cell mediated delayed type hypersensitivity and experimental autoimmunity are suppressed by low doses of heparanase inhibiting non-anticoagulant species of heparin (30).

Eleventh, heparanase activity expressed by platelets, neutrophils and metastatic tumor cells releases active bFGF from ECM and basement membranes. Release of bFGF from storage in ECM may elicit a localized neovascular response in processes such as wound healing, inflammation and tumor development.

Twelfth, among the breakdown products of the ECM generated by heparanase is a tri-sulfated disaccharide that can inhibit T-cell mediated inflammation *in vivo* (31). This inhibition was associated with an inhibitory effect of the disaccharide on the production of biologically active TNFα by activated T cells *in vitro* (31).

Other potential therapeutic applications: Apart from its involvement in tumor cell metastasis, inflammation and autoimmunity, mammalian heparanase may be applied to modulate: bioavailability of heparin-binding growth factors (15); cellular responses to heparin-binding

viral and some bacterial and protozoa infections (33, 33a, 33b); and disintegration of amyloid plaques (34). Heparanase may thus prove useful for conditions such as wound healing, angiogenesis, restenosis, atherosclerosis, inflammation, neurodegenerative diseases and viral infections. Mammalian heparanase can be used to neutralize plasma heparin, as a potential replacement of protamine. Anti-heparanase antibodies may be applied for immunodetection and diagnosis of micrometastases, autoimmune lesions and renal failure in biopsy specimens, plasma samples, and body fluids. Common use in basic research is expected.

The identification of the *hpa* gene encoding for heparanase enzyme will enable the production of a recombinant enzyme in heterologous expression systems. Availability of the recombinant protein will pave the way for solving the protein structure function relationship and will provide a tool for developing new inhibitors.

Viral Infection: The presence of heparan sulfate on cell surfaces have been shown to be the principal requirement for the binding of Herpes Simplex (33) and Dengue (33a) viruses to cells and for subsequent infection of the cells. Removal of the cell surface heparan sulfate by heparanase may therefore abolish virus infection. In fact, treatment of cells with bacterial

rendered the cells at least partially resistant to virus infection (33). There are some indications that the cell surface heparan sulfate is also involved in HIV infection (33b).

Neurodegenerative diseases: Heparan sulfate proteoglycans were identified in the prion protein amyloid plaques of Genstmann-Straussler Syndrome, Creutzfeldt-Jakob disease and Scrape (34). Heparanase may disintegrate these amyloid plaques which are also thought to play a role in the pathogenesis of Alzheimer's disease.

Restenosis and Atherosclerosis: Proliferation of arterial smooth muscle cells (SMCs) in response to endothelial injury and accumulation of cholesterol rich lipoproteins are basic events in the pathogenesis of atherosclerosis and restenosis (35). Apart from its involvement in SMC proliferation (i.e., low affinity receptors for heparin-binding growth factors), HS is also involved in lipoprotein binding, retention and uptake (36). It was demonstrated that HSPG and lipoprotein lipase participate in a novel catabolic pathway that may allow substantial cellular and interstitial accumulation of cholesterol rich lipoproteins (32). The latter pathway is expected to be highly atherogenic by promoting accumulation of apoB and apoE rich lipoproteins (i.e. LDL, VLDL, chylomicrons), independent of feed back inhibition by the cellular sterol content. Removal of SMC HS by

accumulation and thus may halt the progression of restenosis and atherosclerosis.

Gene therapy:

The ultimate goal in the management of inherited as well as acquired diseases is a rational therapy with the aim to eliminate the underlying biochemical defects associated with the disease rather then symptomatic treatment. Gene therapy is a promising candidate to meet these objectives. Initially it was developed for treatment of genetic disorders, however, the consensus view today is that it offers the prospect of providing therapy for a variety of acquired diseases, including cancer, viral infections, vascular diseases and neurodegenerative disorders.

The gene-based therapeutic can act either intracellularly, affecting only the cells to which it is delivered, or extracellularly, using the recipient cells as local endogenous factories for the therapeutic product(s). The application of gene therapy may follow any of the following strategies: (i) prophylactic gene therapy, such as using gene transfer to protect cells against viral infection; (ii) cytotoxic gene therapy, such as cancer therapy, where genes encode cytotoxic products to render the target cells vulnerable to attack by the normal immune response; (iii) biochemical correction, primarily for the treatment of single gene defects, where a normal copy of

To allow efficient transfer of the therapeutic genes, a variety of gene delivery techniques have been developed based on viral and non-viral vector systems. The most widely used and most efficient systems for delivering genetic material into target cells are viral vectors. So far, 329 clinical studies (phase I, I/II and II) with over 2,500 patients have been initiated Worldwide since 1989 (50).

The approach of gene addition pose serious barriers. The expression of many genes is tightly regulated and context dependent, so achieving the correct balance and function of expression is challenging. The gene itself is often quite large, containing many exons and introns. The delivery vector is usually a virus, which can infect with a high efficiency but may, on the other hand, induce immunological response and consequently decreases effectiveness, especially upon secondary administration. Most of the current expression vector-based gene therapy protocols fail to achieve clinically significant transgene expression required for treating genetic diseases. Apparently, it is difficult to deliver enough virus to the right cell type to elicit an effective and therapeutic effect (51).

Homologous recombination, which was initially considered to be of limited use for gene therapy because of its low frequency in mammalian cells, has recently emerged as a potential strategy for developing gene

These studies aimed at either gene disruption or gene correction and include RNA DNA chimeric oligonucleotides, small or large homologous DNA fragments, or adeno-associated viral vectors. Most of these studies show a reasonable frequency of homologous recombination, which warrants further *in vivo* testing (52). Homologous recombination-based gene therapy has the potential to develop into a powerful therapeutic modality for genetic diseases. It can offer permanent expression and normal regulation of corrected genes in appropriate cells or organs and probably can be used for treating dominantly inherited diseases such as polycystic kidney disease.

Genomic sequences function in regulation of gene expression:

The efficient expression of therapeutic genes in target cells or tissues is an important component of efficient and safe gene therapy. The expression of genes is driven by the promoter region upstream of the coding sequence, although regulation of expression may be supplemented by farther upstream or downstream DNA sequences or DNA in the introns of the gene. Since this important information is embedded in the DNA, the description of gene structure is crucial to the analysis of gene regulation. Characterization of cell specific or tissue specific promoters, as well as other tissue specific regulatory elements enables the use of such sequences to direct efficient cell specific, or developmental stage specific gene

drugs. Identification of transcription factors and other regulatory proteins required for proper gene expression will point at new potential targets for modulating gene expression, when so desired or required.

Efficient expression of many mammalian genes depends on the presence of at least one intron. The expression of mouse thymidylate synthase (TS) gene, for example, is greatly influenced by intron sequences. The addition of almost any of the introns from the mouse TS gene to an intronless TS minigene leads to a large increase in expression (42). The involvement of intron 1 in the regulation of expression was demonstrated for many other genes. In human factor IX (hFIX), intron 1 is able to increase the expression level about 3 fold mare as compared to that of the hFIX cDNA (43). The expression enhancing activity of intron 1 is due to efficient functional splicing sequences, present in the precursor mRNA. By being efficiently assembled into spliceosome complexes, transcripts with splicing sequences may be better protected in the nucleus from random degradations, than those without such sequences (44).

A forward-inserted intron1-carrying hFIX expression cassette suggested to be useful for directed gene transfer, while for retroviral-mediated gene transfer system, reversely-inserted intron 1-carrying hFIX expression cassette was considered (43).

Ki-ras genes of human, mouse and rat. This cis-acting regulatory sequence confers strong transcription enhancer activity that is differentially modulated by steroid hormones in metastatic and nonmetastatic subpopulations. Perturbations in the regulatory activities of such cis-acting sequences may play an important role in governing oncogenic potency of Ha-ras through transcriptional control mechanisms (45).

Intron sequences affect tissue specific, as well as inducible gene expression. A 182 bp intron 1 DNA segment of the mouse Col2a1 gene contains the necessary information to confer high-level, temporally correct, chondrocyte expression on a reporter gene in intact mouse embryos, while Col2a1 promoter sequences are dispensable for chondrocyte expression (46). In Col1A1 gene the intron plays little or no role in constitutive expression of collagen in the skin, and in cultured cells derived from the skin, however, in the lungs of young mice, intron deletion results in decrease of expression to less than 50 % (47).

A classical enhancer activity was shown in the 2 kb intron fragment in bovine beta-casein gene. The enhancer activity was largely dependent on the lactogenic hormones, especially prolactin. It was suggested that several elements in the intron-1 of the bovine beta-casein gene cooperatively interact not only with each other but also with its promoter for hormonal

Identification and characterization of regulatory elements in genomic non-coding sequences, such as introns, provides a tool for designing and constructing novel vectors for tissue specific, hormone regulated or any other defined expression pattern, for gene therapy. Such an expression cassette was developed, utilizing regulatory elements from the human cytokeratin 18 (K18) gene, including 5' genomic sequences and one of its introns. This cassette efficiently expresses reporter genes, as well as the human cystic fibrosis transmembrane conductance regulator (CFTR) gene, in cultured lung epithelial cells (49).

Alternative splicing:

Alternative splicing of pre mRNA is a powerful and versatile regulatory mechanism that can effect quantitative control of gene expression and functional diversification of proteins. It contributes to major developmental decisions and also to a fine-tuning of gene function. Genetic and biochemical approaches have identified cis-acting regulatory elements and trans-acting factors that control alternative splicing of specific mRNAs. This mechanism results in the generation of variant isoforms of various proteins from a single gene. These include cell surface molecules such as CD44, receptors, cytokines such as VEGF and enzymes. Products of alternatively spliced transcripts differ in their expression pattern, substrate

The FGF receptor RNA undergoes alternative splicing which results in the production of several isoforms, which exhibit different ligand binding specificities. The alternative splicing is regulated in a cell specific manner (53).

Alternative spliced mRNAs are often correlated with malignancy. An increase in specific splice variant of tyrosinase was identified in murine melanomas (54). Multiple splicing variants of estrogen receptor are present in individual human breast tumors. CD44 has various isoform, some are characteristic of malignant tissues.

Identification of tumor specific alternative splice variants provide new tool for cancer diagnostics. CD44 variants have been used for detection of malignancy in urine samples from patients with urothelial cancer by competitive RT-PCR (55). CD44 exon 6 was suggested as prognostic indicator of metastasis in breast cancer (56).

Different enzymes or polypeptides generated by alternative splicing may have different function or catalytic specificity. The identification and characterization of the enzyme forms, which are involved in pathological processes, is crucial for the design of appropriate and efficient drugs.

Modulation of gene expression – Antisense technology:

An antisense oligonucleotide (e.g., antisense

(64). According to the Watson-Crick base pairing, heterocyclic bases of the antisense oligonucleotide form hydrogen bonds with the heterocyclic bases of target single-stranded nucleic acids (RNA or single-stranded DNA), whereas according to the Hoogsteen base pairing, the heterocyclic bases of the target nucleic acid are double-stranded DNA, wherein a third strand is accommodated in the major groove of the B-form DNA duplex by Hoogsteen and anti-Hoogsteen base pairing to form a triple helix structure.

According to both the Watson-Crick and the Hoogsteen base pairing models, antisense oligonucleotides have the potential to regulate gene expression and to disrupt the essential functions of the nucleic acids in cells. Therefore, antisense oligonucleotides have possible uses in modulating a wide range of diseases in which gene expression is altered.

Since the development of effective methods for chemically synthesizing oligonucleotides, these molecules have been extensively used in biochemistry and biological research and have the potential use in medicine, since carefully devised oligonucleotides can be used to control gene expression by regulating levels of transcription, transcripts and or translation.

Oligodeoxyribonucleotides as long as 100 base pairs (bp) are routinely synthesized by solid phase methods using commercially available,

also much less stable than oligodeoxyribonucleotides, a fact which has contributed to the more prevalent use of oligodeoxyribonucleotides in medical and biological research, directed at, for example, the regulation of transcription or translation levels.

Gene expression involves few distinct and well regulated steps. The first major step of gene expression involves transcription of a messenger RNA (mRNA) which is an RNA sequence complementary to the antisense (i.e., -) DNA strand, or, in other words, identical in sequence to the DNA sense (i.e., +) strand, composing the gene. In eukaryotes, transcription occurs in the cell nucleus.

The second major step of gene expression involves translation of a protein (e.g., enzymes, structural proteins, secreted proteins, gene expression factors, etc.) in which the mRNA interacts with ribosomal RNA complexes (ribosomes) and amino acid activated transfer RNAs (tRNAs) to direct the synthesis of the protein coded for by the mRNA sequence.

Initiation of transcription requires specific recognition of a promoter DNA sequence located upstream to the coding sequence of a gene by an RNA-synthesizing enzyme -- RNA polymerase. This recognition is preceded by sequence-specific binding of one or more transcription factors to the promoter sequence. Additional proteins which bind at or close to the

promoter, but whose binding prohibits the action of RNA polymerase, are known as repressors.

There are also evidence that in some cases gene expression is downregulated by endogenous antisense RNA repressors that bind a complementary mRNA transcript and thereby prevent its translation into a functional protein.

Thus, gene expression is typically upregulated by transcription factors and enhancers and downregulated by repressors.

However, in many disease situation gene expression is impaired. In many cases, such as different types of cancer, for various reasons the expression of a specific endogenous or exogenous (e.g., of a pathogen such as a virus) gene is upregulated. Furthermore, in infectious diseases caused by pathogens such as parasites, bacteria or viruses, the disease progression depends on expression of the pathogen genes, this phenomenon may also be considered as far as the patient is concerned as upregulation of exogenous genes.

Most conventional drugs function by interaction with and modulation of one or more targeted endogenous or exogenous proteins, e.g., enzymes. Such drugs, however, typically are not specific for targeted proteins but interact with other proteins as well. Thus, a relatively large dose of drug

Typical daily doses of drugs are from 10^{-5} - 10^{-1} millimoles per kilogram of body weight or 10^{-3} - 10 millimoles for a 100 kilogram person. If this modulation instead could be effected by interaction with and inactivation of mRNA, a dramatic reduction in the necessary amount of drug could likely be achieved, along with a corresponding reduction in side effects. Further reductions could be effected if such interaction could be rendered site-specific. Given that a functioning gene continually produces mRNA, it would thus be even more advantageous if gene transcription could be arrested in its entirety.

Given these facts, it would be advantageous if gene expression could be arrested or downmodulated at the transcription level.

The ability of chemically synthesizing oligonucleotides and analogs thereof having a selected predetermined sequence offers means for downmodulating gene expression. Three types of gene expression modulation strategies may be considered.

At the transcription level, antisense or sense oligonucleotides or analogs that bind to the genomic DNA by strand displacement or the formation of a triple helix, may prevent transcription (64).

At the transcript level, antisense oligonucleotides or analogs that bind target mRNA molecules lead to the enzymatic cleavage of the hybrid

hybrid recognized and destroyed by the RNase H enzyme. Alternatively, such hybrid formation may lead to interference with correct splicing (66). As a result, in both cases, the number of the target mRNA intact transcripts ready for translation is reduced or eliminated.

At the translation level, antisense oligonucleotides or analogs that bind target mRNA molecules prevent, by steric hindrance, binding of essential translation factors (ribosomes), to the target mRNA, a phenomenon known in the art as hybridization arrest, disabling the translation of such mRNAs (67).

Thus, antisense sequences, which as described hereinabove may arrest the expression of any endogenous and/or exogenous gene depending on their specific sequence, attracted much attention by scientists and pharmacologists who were devoted at developing the antisense approach into a new pharmacological tool (68).

For example, several antisense oligonucleotides have been shown to arrest hematopoietic cell proliferation (69), growth (70), entry into the S phase of the cell cycle (71), reduced survival (72) and prevent receptor mediated responses (73). For use of antisense oligonucleotides as antiviral agents the reader is referred to reference 74.

For efficient *in vivo* inhibition of gene expression using antisense

sequence; (ii) solubility in water; (iii) stability against intra- and extracellular nucleases; (iv) capability of penetration through the cell membrane; and (v) when used to treat an organism, low toxicity.

Unmodified oligonucleotides are impractical for use as antisense sequences since they have short *in vivo* half-lives, during which they are degraded rapidly by nucleases. Furthermore, they are difficult to prepare in more than milligram quantities. In addition, such oligonucleotides are poor cell membrane penetraters (75).

Thus it is apparent that in order to meet all the above listed requirements, oligonucleotide analogs need to be devised in a suitable manner. Therefore, an extensive search for modified oligonucleotides has been initiated.

For example, problems arising in connection with double-stranded DNA (dsDNA) recognition through triple helix formation have been diminished by a clever "switch back" chemical linking, whereby a sequence of polypurine on one strand is recognized, and by "switching back", a homopurine sequence on the other strand can be recognized. Also, good helix formation has been obtained by using artificial bases, thereby improving binding conditions with regard to ionic strength and pH.

In addition, in order to improve half-life as well as membrane

Oligonucleotides can be modified either in the base, the sugar or the phosphate moiety. These modifications include, for example, the use of methylphosphonates, monothiophosphates, dithiophosphates, phosphoramidates, phosphate esters, bridged phosphorothioates, bridged phosphoramidates, bridged methylenephosphonates, dephospho internucleotide analogs with siloxane bridges, carbonate bridges, carboxymethyl ester bridges, carbonate bridges, carboxymethyl ester bridges, acetamide bridges, carbamate bridges, thioether bridges, sulfoxy bridges, sulfono bridges, various "plastic" DNAs, α-anomeric bridges and borane derivatives. For further details the reader is referred to reference 76.

International patent application WO 89/12060 discloses various building blocks for synthesizing oligonucleotide analogs, as well as oligonucleotide analogs formed by joining such building blocks in a defined sequence. The building blocks may be either "rigid" (i.e., containing a ring structure) or "flexible" (i.e., lacking a ring structure). In both cases, the building blocks contain a hydroxy group and a mercapto group, through which the building blocks are said to join to form oligonucleotide analogs. The linking moiety in the oligonucleotide analogs is selected from the group consisting of sulfide (-S-), sulfoxide (-SO-), and sulfone ($-SO_2-$). However, the application provides no data supporting the specific binding of an

International patent application WO 92/20702 describe an acyclic oligonucleotide which includes a peptide backbone on which any selected chemical nucleobases or analogs are stringed and serve as coding characters as they do in natural DNA or RNA. These new compounds, known as peptide nucleic acids (PNAs), are not only more stable in cells than their natural counterparts, but also bind natural DNA and RNA 50 to 100 times more tightly than the natural nucleic acids cling to each other (77). PNA oligomers can be synthesized from the four protected monomers containing thymine, cytosine, adenine and guanine by Merrifield solid-phase peptide synthesis. In order to increase solubility in water and to prevent aggregation, a lysine amide group is placed at the C-terminal.

Thus, antisense technology requires pairing of messenger RNA with an oligonucleotide to form a double helix that inhibits translation. The concept of antisense-mediated gene therapy was already introduced in 1978 for cancer therapy. This approach was based on certain genes that are crucial in cell division and growth of cancer cells. Synthetic fragments of genetic substance DNA can achieve this goal. Such molecules bind to the targeted gene molecules in RNA of tumor cells, thereby inhibiting the translation of the genes and resulting in dysfunctional growth of these cells. Other mechanisms has also been proposed. These strategies have been

are typically synthesized in lengths of 13-30 nucleotides. The life span of oligonucleotide molecules in blood is rather short. Thus, they have to be chemically modified to prevent destruction by ubiquitous nucleases present in the body. Phosphorothioates are very widely used modification in antisense oligonucleotide ongoing clinical trials (57). A new generation of antisense molecules consist of hybrid antisense oligonucleotide with a central portion of synthetic DNA while four bases on each end have been modified with 2'O methyl ribose to resemble RNA. In preclinical studies in laboratory animals, such compounds have demonstrated greater stability to metabolism in body tissues and an improved safety profile when compared with the first-generation unmodified phosphorothioate (Hybridon Inc. news). Dosens of other nucleotide analogs have also been tested in antisense technology.

RNA oligonucleotides may also be used for antisense inhibition as they form a stable RNA-RNA duplex with the target, suggesting efficient inhibition. However, due to their low stability RNA oligonucleotides are typically expressed inside the cells using vectors designed for this purpose. This approach is favored when attempting to target a mRNA that encodes an abundant and long-lived protein (57).

Recent scientific publications have validated the efficacy of antisense

approved by the FDA. This drug Fomivirsen, developed by Isis, is indicated for local treatment of cytomegalovirus in patients with AIDS who are intolerant of or have a contraindication to other treatments for CMV retinitis or who were insufficiently responsive to previous treatments for CMV retinitis (Pharmacotherapy News Network).

Several antisense compounds are now in clinical trials in the United States. These include locally administered antivirals, systemic cancer therapeutics. Antisense therapeutics has the potential to treat many life-threatening diseases with a number of advantages over traditional drugs. Traditional drugs intervene after a disease-causing protein is formed. Antisense therapeutics, however, block mRNA transcription/translation and intervene before a protein is formed, and since antisense therapeutics target only one specific mRNA, they should be more effective with fewer side effects than current protein-inhibiting therapy.

A second option for disrupting gene expression at the level of transcription uses synthetic oligonucleotides capable of hybridizing with double stranded DNA. A triple helix is formed. Such oligonucleotides may prevent binding of transcription factors to the gene's promoter and therefore inhibit transcription. Alternatively, they may prevent duplex unwinding and, therefore, transcription of genes within the triple helical structure.

DNA sequences it is possible to synthesize oligonucleotides that will effectively compete with the native DNA sequences for available transcription factors *in vivo*. This approach requires the identification of gene specific transcription factor (57).

Indirect inhibition of gene expression was demonstrated for matrix metalloproteinase genes (MMP-1, -3, and -9), which are associated with invasive potential of human cancer cells. E1AF is a transcription activator of MMP genes. Expression of E1AF antisense RNA in HSC3AS cells showed decrease in mRNA and protein levels of MMP-1, -3, and -9. Moreover, HSC3AS showed lower invasive potential in vitro and *in vivo*. These results imply that transfection of antisense inhibits tumor invasion by down-regulating MMP genes (58).

Ribozymes:

Ribozymes are being increasingly used for the sequence-specific inhibition of gene expression by the cleavage of mRNAs encoding proteins of interest. The possibility of designing ribozymes to cleave any specific target RNA has rendered them valuable tools in both basic research and therapeutic applications. In the therapeutics area, ribozymes have been exploited to target viral RNAs in infectious diseases, dominant oncogenes in cancers and specific somatic mutations in genetic disorders. Most

transgenic animal research, gene target validation and pathway elucidation. Several ribozymes are in various stages of clinical trials. ANGIOZYME was the first chemically synthesized ribozyme to be studied in human clinical trials. ANGIOZYME specifically inhibits formation of the VEGF-r (Vascular Endothelial Growth Factor receptor), a key component in the angiogenesis pathway. Ribozyme Pharmaceuticals, Inc., as well as other firms have demonstrated the importance of anti-angiogenesis therapeutics in animal models. HEPTAZYME, a ribozyme designed to selectively destroy Hepatitis C Virus (HCV) RNA, was found effective in decreasing Hepatitis C viral RNA in cell culture assays (Ribozyme Pharmaceuticals, Incorporated - WEB home page).

Gene disruption in animal models:

The emergence of gene inactivation by homologous recombination methodology in embryonic stem cells has revolutionized the field of mouse genetics. The availability of a rapidly growing number of mouse null mutants has represented an invaluable source of knowledge on mammalian development, cellular biology and physiology, and has provided many models for human inherited diseases. Animal models are required for an effective drug delivery development program and evaluation of gene therapy approach. The improvement of the original knockout strategy, as

manipulations that can be produced. Additional methods have been developed to provide versatile research tools: Double replacement method, sequential gene targeting, conditional cell type specific gene targeting, single copy integration method, inducible gene targeting, gene disruption by viral delivery, replacing one gene with another, the so called knock-in method and the induction of specific balanced chromosomal translocation. It is now possible to introduce a point mutation as a unique change in the entire genome, therefore allowing very fine dissection of gene function *in vivo*. Furthermore, the advent of methods allowing conditional gene targeting opens the way for analysis of consequence of a particular mutation in a defined organ and at a specific time during the life of the experimental animal (59).

DNA vaccination:

Observations in the early 1990s that plasmid DNA could directly transfect animal cells *in vivo* sparked exploration of the use of DNA plasmids to induce immune response by direct injection into animal of DNA encoding antigenic protein. When a DNA vaccine plasmid enters the eukaryotic cell, the protein it encodes is transcribed and translated within the cell. In the case of pathogens, these proteins are presented to the immune system in their native form, mimicking the presentation of antigens

infectious diseases, as well as for allergy and for cancer. The central hypothesis behind active specific immunotherapy for cancer is that tumor cells express unique antigens that should stimulate the immune system. The first DNA vaccine against tumor was carcino-embrionic antigen (CEA). DNA vaccinated animals expressed immunoprotection and immunotherapy of human CEA-expressing syngeneic mouse colon and breast carcinoma (61). In a mouse model of neuroblastoma, DNA immunization with HuD resulted in tumor growth inhibition with no neurological disease (60). Immunity to the brown locus protein, gp^{75} tyrosinase-related protein-1, associated with melanoma, was investigated in a syngeneic mouse model. Priming with human gp75 DNA broke tolerance to mouse gp75. Immunity against mouse gp75 provided significant tumor protection (60).

Glycosyl hydrolases:

Glycosyl hydrolases are a widespread group of enzymes that hydrolyze the o-glycosidic bond between two or more carbohydrates or between a carbohydrate and a noncarbohydrate moiety. The enzymatic hydrolysis of glycosidic bond occurs by using major one or two mechanisms leading to overall retention or inversion of the anomeric configuration. In both mechanisms catalysis involves two residues: a proton donor and a nucleophile. Glycosyl hydrolyses have been classified into 58 families

they all hydrolyze the glycosidic bond in a general acid catalysis mechanism, with retention of the anomeric configuration. The mechanism involves two glutamic acid residues, which are the proton donors and the nucleophile, with an aspargine always preceding the proton donor. Analyses of a set of known 3D structures from this group revealed that their catalytic domains, despite the low level of sequence identity, adopt a similar (α/β) 8 fold with the proton donor and the nucleophile located at the C-terminal ends of strands β4 and β7, respectively. Mutations in the functional conserved amino acids of lysosomal glycosyl hydrolases were identified in lysosomal storage diseases.

Lysosomal glycosyl hydrolases including β-glucuronidase, β-manosidase, β-glucocerebrosidase, β-galactosidase and α-L iduronidase, are all exo-glycosyl hydrolases, belong to the GH-A clan and share a similar catalytic site. However, many endo-glucanases from various organisms, such as bacterial and fungal xylenases and cellulases share this catalytic domain.

Genomic sequence of hpa gene and its implications:

It is well established that heparanase activity is correlated with cancer metastasis. This correlation was demonstrated at the level of enzymatic activity as well as the levels of protein and *hpa* cDNA expression

by several means. The genomic region, encoding the *hpa* gene and the surrounding, provides a new powerful tool for regulation of heparanase activity at the level of gene expression. Regulatory sequences may reside in noncoding regions both upstream and downstream the transcribed region as well as in intron sequences. A DNA sequence upstream of the transcription start site contains the promoter region and potential regulatory elements. Regulatory factors, which interact with the promoter region may be identified and be used as potential drugs for inhibition of cancer, metastasis and inflammation. The promoter region can be used to screen for inhibitors of heparanase gene expression. Furthermore, the *hpa* promoter can be used to direct cell specific, particularly cancer cell specific, expression of foreign genes, such as cytotoxic or apoptotic genes, in order to specifically destroy cancer cells.

Cancer and yet unknown related genetic disorders may involve rearrangements and mutations in the heparanase gene, either in coding or non-coding regions. Such mutations may affect expression level or enzymatic activity. The genomic sequence of *hpa* enables the amplification of specific genomic DNA fragments, identification and diagnosis of mutations.

There is thus a widely recognized need for, and it would be highly

encoding a polypeptide having heparanase activity, vector including same,

genetically modified cells expressing heparanase and a recombinant protein having heparanase activity, as well as antisense oligonucleotides, constructs and ribozymes which can be used for down regulation heparanase activity.

SUMMARY OF THE INVENTION

Cloning of the human *hpa* gene which encodes heparanase, and expression of recombinant heparanase by transfected host cells is reported herein, as well as downregulation of heparanase activity by antisense technology.

A purified preparation of heparanase isolated from human hepatoma cells was subjected to tryptic digestion and microsequencing. The YGPDVGQPR (SEQ ID NO:8) sequence revealed was used to screen EST databases for homology to the corresponding back translated DNA sequence. Two closely related EST sequences were identified and were thereafter found to be identical. Both clones contained an insert of 1020 bp which included an open reading frame of 973 bp followed by a 27 bp of 3' untranslated region and a Poly A tail. Translation start site was not identified.

Cloning of the missing 5' end of *hpa* was performed by PCR amplification of DNA from placenta Marathon RACE cDNA composite

identified 3' encoding EST clones was obtained. The joined cDNA fragment (*hpa*), 1721 bp long (SEQ ID NO:9), contained an open reading frame which encodes a polypeptide of 543 amino acids (SEQ ID NO:10) with a calculated molecular weight of 61,192 daltons.

Cloning an extended 5' sequence was enabled from the human SK-hep1 cell line by PCR amplification using the Marathon RACE. The 5' extended sequence of the SK-hep1 *hpa* cDNA was assembled with the sequence of the *hpa* cDNA isolated from human placenta (SEQ ID NO:9). The assembled sequence contained an open reading frame, SEQ ID NOs: 13 and 15, which encodes, as shown in SEQ ID NOs:14 and 15, a polypeptide of 592 amino acids with a calculated molecular weight of 66,407 daltons.

The ability of the *hpa* gene product to catalyze degradation of heparan sulfate in an *in vitro* assay was examined by expressing the entire open reading frame of *hpa* in insect cells, using the Baculovirus expression system. Extracts and conditioned media of cells infected with virus containing the *hpa* gene, demonstrated a high level of heparan sulfate degradation activity both towards soluble ECM-derived HSPG and intact ECM. This degradation activity was inhibited by heparin, which is another substrate of heparanase. Cells infected with a similar construct containing no *hpa* gene had no such activity, nor did non-infected cells. The ability of

[illegible] a [illegible] [illegible]

The expression pattern of *hpa* RNA in various tissues and cell lines was investigated using RT-PCR. It was found to be expressed only in tissues and cells previously known to have heparanase activity.

A panel of monochromosomal human/CHO and human/mouse somatic cell hybrids was used to localize the human heparanase gene to human chromosome 4. The newly isolated heparanase sequence can be used to identify a chromosome region harboring a human heparanase gene in a chromosome spread.

A human genomic library was screened and the human locus harboring the heparanase gene isolated, sequenced and characterized. Alternatively spliced heparanase mRNAs were identified and characterized. The human heparanase promoter has been isolated, identified and positively tested for activity. The mouse heparanase promoter has been isolated and identified as well. Antisense heparanase constructs were prepared and their influence on cells *in vitro* tested. A predicted heparanase active site was identified. And finally, the presence of sequences hybridizing with human heparanase sequences was demonstrated for a variety of mammalians and for an avian.

According to one aspect of the present invention there is provided an isolated nucleic acid comprising a genomic, complementary or composite

According to further features in preferred embodiments of the invention described below, the polynucleotide or a portion thereof is hybridizable with SEQ ID NOs: 9, 13, 42, 43 or a portion thereof at 68 °C in 6 x SSC, 1 % SDS, 5 x Denharts, 10 % dextran sulfate, 100 µg/ml salmon sperm DNA, and ^{32}p labeled probe and wash at 68 °C with 3 x SSC and 0.1 % SDS.

According to still further features in the described preferred embodiments the polynucleotide or a portion thereof is at least 60 % identical with SEQ ID NOs: 9, 13, 42, 43 or portions thereof as determined using the Bestfit procedure of the DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the university of Wisconsin (gap creation penalty - 12, gap extension penalty - 4).

According to still further features in the described preferred embodiments the polypeptide is as set forth in SEQ ID NOs:10, 14, 44 or portions thereof.

According to still further features in the described preferred embodiments the polypeptide is at least 60 % homologous to SEQ ID NOs:10, 14, 44 or portions thereof as determined with the Smith-Waterman algorithm, using the Bioaccelerator platform developed by Compugene (gapop: 10.0, gapext: 0.5, matrix: blosum62).

According to additional aspects of the present invention there are provided a nucleic acid construct (vector) comprising the isolated nucleic acid described herein and a host cell comprising the construct.

According to a further aspect of the present invention there is provided an antisense oligonucleotide comprising a polynucleotide or a polynucleotide analog of at least 10 bases being hybridizable *in vivo*, under physiological conditions, with a portion of a polynucleotide strand encoding a polypeptide having heparanase catalytic activity.

According to an additional aspect of the present invention there is provided a method of *in vivo* downregulating heparanase activity comprising the step of *in vivo* administering the antisense oligonucleotide herein described.

According to yet an additional aspect of the present invention there is provided a pharmaceutical composition comprising the antisense oligonucleotide herein described and a pharmaceutically acceptable carrier.

According to still an additional aspect of the present invention there is provided a ribozyme comprising the antisense oligonucleotide described herein and a ribozyme sequence.

According to a further aspect of the present invention there is provided an antisense nucleic acid construct comprising a promoter

under physiological conditions, with a portion of a polynucleotide strand encoding a polypeptide having heparanase catalytic activity.

According to further features in preferred embodiments of the invention described below, the polynucleotide strand encoding the polypeptide having heparanase catalytic activity is as set forth in SEQ ID NOs: 9, 13, 42 or 43.

According to still further features in the described preferred embodiments the polypeptide having heparanase catalytic activity is as set forth in SEQ ID NOs: 10, 14 or 44.

According to still a further aspect of the present invention there is provided a method of *in vivo* downregulating heparanase activity comprising the step of *in vivo* administering the antisense nucleic acid construct herein described.

According to yet a further aspect of the present invention there is provided a pharmaceutical composition comprising the antisense nucleic acid construct herein described and a pharmaceutically acceptable carrier.

According to a further aspect of the present invention there is provided a nucleic acid construct comprising a polynucleotide sequence functioning as a promoter, the polynucleotide sequence is derived from SEQ ID NO:42 and includes at least nucleotides 2535-2635 thereof or from SEQ

According to a further aspect of the present invention there is provided a method of expressing a polynucleotide sequence comprising the step of ligating the polynucleotide sequence into the nucleic acid construct described above, downstream of the polynucleotide sequence derived from SEQ ID NOs:42 or 43.

According to a further aspect of the present invention there is provided a recombinant protein comprising a polypeptide having heparanase catalytic activity.

According to further features in preferred embodiments of the invention described below, the polypeptide includes at least a portion of SEQ ID NOs:10, 14 or 44.

According to still further features in the described preferred embodiments the protein is encoded by a polynucleotide hybridizable with SEQ ID NOs: 9, 13, 42, 43 or a portion thereof at 68 °C in 6 x SSC, 1 % SDS, 5 x Denharts, 10 % dextran sulfate, 100 µg/ml salmon sperm DNA, and ^{32}p labeled probe and wash at 68 °C with 3 x SSC and 0.1 % SDS.

According to still further features in the described preferred embodiments the protein is encoded by a polynucleotide at least 60 % identical with SEQ ID NOs: 9, 13, 42, 43 or portions thereof as determined using the Bestfit procedure of the DNA sequence analysis software package

According to a further aspect of the present invention there is provided a pharmaceutical composition comprising, as an active ingredient, the recombinant protein herein described.

According to a further aspect of the present invention there is provided a method of identifying a chromosome region harboring a heparanase gene in a chromosome spread comprising the steps of (a) hybridizing the chromosome spread with a tagged polynucleotide probe encoding heparanase; (b) washing the chromosome spread, thereby removing excess of non-hybridized probe; and (c) searching for signals associated with the hybridized tagged polynucleotide probe, wherein detected signals being indicative of a chromosome region harboring a heparanase gene.

According to a further aspect of the present invention there is provided a method of *in vivo* eliciting anti-heparanase antibodies comprising the steps of administering a nucleic acid construct including a polynucleotide segment corresponding to at least a portion of SEQ ID NOs:9, 13 or 43 and a promoter for directing the expression of said polynucleotide segment *in vivo*. Accordingly, there is provided also a DNA vaccine for *in vivo* eliciting anti-heparanase antibodies comprising a nucleic acid construct including a polynucleotide segment corresponding to at least

The present invention can be used to develop new drugs to inhibit tumor cell metastasis, inflammation and autoimmunity. The identification of the *hpa* gene encoding for heparanase enzyme enables the production of a recombinant enzyme in heterologous expression systems. Additional features, advantages, uses and applications of the present invention in biological science and in diagnostic and therapeutic medicine are described hereinafter.

BRIEF DESCRIPTION OF THE DRAWINGS

The invention herein described, by way of example only, with reference to the accompanying drawings, wherein:

FIG. 1 presents nucleotide sequence and deduced amino acid sequence of *hpa* cDNA. A single nucleotide difference at position 799 (A to T) between the EST (Expressed Sequence Tag) and the PCR amplified cDNA (reverse transcribed RNA) and the resulting amino acid substitution (Tyr to Phe) are indicated above and below the substituted unit, respectively. Cysteine residues and the poly adenylation consensus sequence are underlined. The asterisk denotes the stop codon TGA.

FIG. 2 demonstrates degradation of soluble sulfate labeled HSPG substrate by lysates of High Five cells infected with pF*hpa*2 virus. Lysates

soluble HSPG (peak I). The incubation medium was then subjected to gel filtration on Sepharose 6B. Low molecular weight HS degradation fragments (peak II) were produced only during incubation with the pF*hpa*2 infected cells, but there was no degradation of the HSPG substrate (◇) by lysates of pF2 infected cells.

FIGs. 3a-b demonstrate degradation of soluble sulfate labeled HSPG substrate by the culture medium of pF*hpa*2 and pF*hpa*4 infected cells. Culture media of High Five cells infected with pF*hpa*2 (3a) or pF*hpa*4 (3b) viruses (●), or with control viruses (□) were incubated (18 h, 37 °C) with sulfate labeled ECM-derived soluble HSPG (peak I, ◇). The incubation media were then subjected to gel filtration on Sepharose 6B. Low molecular weight HS degradation fragments (peak II) were produced only during incubation with the *hpa* gene containing viruses. There was no degradation of the HSPG substrate by the culture medium of cells infected with control viruses.

FIG. 4 presents size fractionation of heparanase activity expressed by pF*hpa*2 infected cells. Culture medium of pF*hpa*2 infected High Five cells was applied onto a 50 kDa cut-off membrane. Heparanase activity (conversion of the peak I substrate, (◇) into peak II HS degradation fragments) was found in the high (> 50 kDa) (●), but not low (< 50 kDa) (○)

FIGs. 5a-b demonstrate the effect of heparin on heparanase activity expressed by pF*hpa*2 and pF*hpa*4 infected High Five cells. Culture media of pF*hpa*2 (5a) and pF*hpa*4 (5b) infected High Five cells were incubated (18 h, 37 °C) with sulfate labeled ECM-derived soluble HSPG (peak I, ◦) in the absence (•) or presence (◦) of 10 μg/ml heparin. Production of low molecular weight HS degradation fragments was completely abolished in the presence of heparin, a potent inhibitor of heparanase activity (6, 7).

FIGs. 6a-b demonstrate degradation of sulfate labeled intact ECM by virus infected High Five and Sf21 cells. High Five (6a) and Sf21 (6b) cells were plated on sulfate labeled ECM and infected (48 h, 28 °C) with pF*hpa*4 (•) or control pF1 (□) viruses. Control non-infected Sf21 cells (R) were plated on the labeled ECM as well. The pH of the cultured medium was adjusted to 6.0 - 6.2 followed by 24 h incubation at 37 °C. Sulfate labeled material released into the incubation medium was analyzed by gel filtration on Sepharose 6B. HS degradation fragments were produced only by cells infected with the *hpa* containing virus.

FIG. 7a-b demonstrate degradation of sulfate labeled intact ECM by virus infected cells. High Five (7a) and Sf21 (7b) cells were plated on sulfate labeled ECM and infected (48 h, 28 °C) with pF*hpa*4 (•) or control pF1 (□) viruses. Control non-infected Sf21 cells (R) were plate on labeled

fragments released into the incubation medium was analyzed by gel filtration on Sepharose 6B. HS degradation fragments were produced only by cells infected with the *hpa* containing virus.

FIGs. 8a-b demonstrate degradation of sulfate labeled intact ECM by the culture medium of pF*hpa4* infected cells. Culture media of High Five (8a) and Sf21 (8b) cells that were infected with pF*hpa*4 (•) or control pF1 (□) viruses were incubated (48 h, 37 °C, pH 6.0) with intact sulfate labeled ECM. The ECM was also incubated with the culture medium of control non-infected Sf21 cells (R). Sulfate labeled material released into the reaction mixture was subjected to gel filtration analysis. Heparanase activity was detected only in the culture medium of pF*hpa*4 infected cells.

FIGs. 9a-b demonstrate the effect of heparin on heparanase activity in the culture medium of pF*hpa*4 infected cells. Sulfate labeled ECM was incubated (24 h, 37 °C, pH 6.0) with culture medium of pF*hpa*4 infected High Five (9a) and Sf21 (9b) cells in the absence (•) or presence (V) of 10 µg/ml heparin. Sulfate labeled material released into the incubation medium was subjected to gel filtration on Sepharose 6B. Heparanase activity (production of peak II HS degradation fragments) was completely inhibited in the presence of heparin.

FIGs. 10a-b demonstrate purification of recombinant heparanase on

fractions was performed with 0.35 - 2 M NaCl gradient (◦). Heparanase activity in the eluted fractions is demonstrated in Figure 10a (•). Fractions 15-28 were subjected to 15 % SDS-polyacrylamide gel electrophoresis followed by silver nitrate staining. A correlation is demonstrated between a major protein band (MW ~ 63,000) in fractions 19 - 24 and heparanase activity.

FIGs. 11a-b demonstrate purification of recombinant heparanase on a Superdex 75 gel filtration column. Active fractions eluted from heparin-Sepharose (Figure 10a) were pooled, concentrated and applied onto Superdex 75 FPLC column. Fractions were collected and aliquots of each fraction were tested for heparanase activity (c, Figure 11a) and analyzed by SDS-polyacrylamide gel electrophoresis followed by silver nitrate staining (Figure 11b). A correlation is seen between the appearance of a major protein band (MW ~ 63,000) in fractions 4 - 7 and heparanase activity.

FIGs. 12a-e demonstrate expression of the *hpa* gene by RT-PCR with total RNA from human embryonal tissues (12a), human extra-embryonal tissues (12b) and cell lines from different origins (12c-e). RT-PCR products using *hpa* specific primers (I), primers for GAPDH housekeeping gene (II), and control reactions without reverse transcriptase demonstrating absence of genomic DNA or other contamination in RNA samples (III). M- DNA

5 - adrenal. For 12b: lane 1 - kidney, lane 2 - placenta (8 weeks), lane 3 - placenta (11 weeks), lanes 4-7 - mole (complete hydatidiform mole), lane 8 - cytotrophoblast cells (freshly isolated), lane 9 - cytotrophoblast cells (1.5 h *in vitro*), lane 10 - cytotrophoblast cells (6 h *in vitro*), lane 11 - cytotrophoblast cells (18 h *in vitro*), lane 12 - cytotrophoblast cells (48 h *in vitro*). For 12c: lane 1 - JAR bladder cell line, lane 2 - NCITT testicular tumor cell line, lane 3 - SW-480 human hepatoma cell line, lane 4 - HTR (cytotrophoblasts transformed by SV40), lane 5 - HPTLP-I hepatocellular carcinoma cell line, lane 6 - EJ-28 bladder carcinoma cell line. For 12d: lane 1 - SK-hep-1 human hepatoma cell line, lane 2 - DAMI human megakaryocytic cell line, lane 3 - DAMI cell line + PMA, lane 4 - CHRF cell line + PMA, lane 5 - CHRF cell line. For 12e: lane 1 - ABAE bovine aortic endothelial cells, lane 2 - 1063 human ovarian cell line, lane 3 - human breast carcinoma MDA435 cell line, lane 4 - human breast carcinoma MDA231 cell line.

FIG. 13 presents a comparison between nucleotide sequences of the human *hpa* and a mouse EST cDNA fragment (SEQ ID NO:12) which is 80 % homologous to the 3' end (starting at nucleotide 1066 of SEQ ID NO:9) of the human *hpa*. The aligned termination codons are underlined.

FIG. 14 demonstrates the chromosomal localization of the *hpa* gene

DNA of hamster, mouse and human of were separated on 0.8 % agarose gel

following amplification with *hpa* specific primers. Lane 1 – Lambda DNA digested with *Bst*EII, lane 2 – no DNA control, lanes 3 – 29, PCR amplification products. Lanes 3-5 – human, mouse and hamster genomic DNA, respectively. Lanes 6-29, human monochromosomal somatic cell hybrids representing chromosomes 1-22 and X and Y, respectively. Lane 30 – Lambda DNA digested with *Bst*EII. An amplification product of approximately 2.8 Kb is observed only in lanes 5 and 9, representing human genomic DNA and DNA derived from cell hybrid carrying human chromosome 4, respectively. These results demonstrate that the *hpa* gene is localized in human chromosome 4.

FIG. 15 demonstrates the genomic exon-intron structure of the human *hpa* locus (top) and the relative positions of the lambda clones used as sequencing templates to sequence the locus (below). The vertical rectangles represent exons (E) and the horizontal lines therebetween represent introns (I), upstream (U) and downstream (D) regions. Continuous lines represent DNA fragments, which were used for sequence analysis. The discontinuous line in lambda 6 represent a region, which overlaps with lambda 8 and hence was not analyzed. The plasmid contains a PCR product, which bridges the gap between L3 and L6.

FIG. 16 presents the nucleotide sequence of the genomic region of

the nucleotide sequence. Two predicted transcription start sites are shown in bold.

FIG. 17 presents an alignment of the amino acid sequences of human heparanase, mouse and partial sequences of rat homologues. The human and the mouse sequences were determined by sequence analysis of the isolated cDNAs. The rat sequence is derived from two different EST clones, which represent two different regions (5' and 3') of the rat *hpa* cDNA. The human sequence and the amino acids in the mouse and rat homologues, which are identical to the human sequence, appear in bold.

FIG. 18 presents a heparanase Zoo blot. Ten micrograms of genomic DNA from various sources were digested with *Eco*RI and separated on 0.7 % agarose – TBE gel. Following electrophoresis, the was gel treated with HCl and than with NaOH and the DNA fragments were downward transferred to a nylon membrane (Hybond N+, Amersham) with 0.4 N NaOH. The membrane was hybridized with a 1.6 Kb DNA probe that contained the entire *hpa* cDNA. Lane order: H – Human; M – Mouse; Rt Rat; P – Pig; Cw – Cow; Hr – Horse; S – Sheep; Rb – Rabbit; D – Dog; Ch Chicken; F – Fish. Size markers (Lambda *Bst*eII) are shown on the left

FIG. 19 demonstrates the secondary structure prediction for heparanase performed using the PHD server – Profile network Prediction

predicted as the proton donor is marked by asterisk and the possible nucleophiles are underlined.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention is of a polynucleotide or nucleic acid, referred to hereinbelow interchangeably as *hpa*, *hpa* cDNA or *hpa* gene or identified by its SEQ ID NOs, encoding a polypeptide having heparanase activity, vectors or nucleic acid constructs including same and which are used for over-expression or antisense inhibition of heparanase, genetically modified cells expressing same, recombinant protein having heparanase activity, antisense oligonucleotides and ribozymes for heparanase modulation, and heparanase promoter sequences which can be used to direct the expression of desired genes.

Before explaining at least one embodiment of the invention in detail, it is to be understood that the invention is not limited in its application to the details of construction and the arrangement of the components set forth in the following description or illustrated in the drawings. The invention is capable of other embodiments or of being practiced or carried out in various ways. Also, it is to be understood that the phraseology and terminology employed herein is for the purpose of description and should not be

Cloning of the human and mouse *hpa* genes, cDNAs and genomic sequence (for human), encoding heparanase and expressing recombinant heparanase by transfected cells is reported herein. These are the first mammalian heparanase genes to be cloned.

A purified preparation of heparanase isolated from human hepatoma cells was subjected to tryptic digestion and microsequencing.

The YGPDVGQPR (SEQ ID NO:8) sequence revealed was used to screen EST databases for homology to the corresponding back translated DNA sequences. Two closely related EST sequences were identified and were thereafter found to be identical.

Both clones contained an insert of 1020 bp which includes an open reading frame of 973 bp followed by a 3' untranslated region of 27 bp and a Poly A tail, whereas a translation start site was not identified.

Cloning of the missing 5' end was performed by PCR amplification of DNA from placenta Marathon RACE cDNA composite using primers selected according to the EST clones sequence and the linkers of the composite.

A 900 bp PCR fragment, partially overlapping with the identified 3' encoding EST clones was obtained. The joined cDNA fragment (*hpa*), 1721 bp long (SEQ ID NO:9), contained an open reading frame which

amino acids (SEQ ID NO:10) with a calculated molecular weight of 61,192 daltons.

A single nucleotide difference at position 799 (A to T) between the EST clones and the PCR amplified cDNA was observed. This difference results in a single amino acid substitution (Tyr to Phe) (Figure 1). Furthermore, the published EST sequences contained an unidentified nucleotide, which following DNA sequencing of both the EST clones was resolved into two nucleotides (G and C at positions 1630 and 1631 in SEQ ID NO:9, respectively).

The ability of the *hpa* gene product to catalyze degradation of heparan sulfate in an *in vitro* assay was examined by expressing the entire open reading frame in insect cells, using the Baculovirus expression system.

Extracts and conditioned media of cells infected with virus containing the *hpa* gene, demonstrated a high level of heparan sulfate degradation activity both towards soluble ECM-derived HSPG and intact ECM, which was inhibited by heparin, while cells infected with a similar construct containing no *hpa* gene had no such activity, nor did non-infected cells.

The expression pattern of hpa RNA in various tissues and cell lines was investigated using RT-PCR. It was found to be expressed only in

Cloning an extended 5' sequence was enabled from the human SK-hep1 cell line by PCR amplification using the Marathon RACE. The 5' extended sequence of the SK-hep1 *hpa* cDNA was assembled with the sequence of the *hpa* cDNA isolated from human placenta (SEQ ID NO:9). The assembled sequence contained an open reading frame, SEQ ID NOs: 13 and 15, which encodes, as shown in SEQ ID NOs:14 and 15, a polypeptide of 592 amino acids, with a calculated molecular weight of 66,407 daltons. This open reading frame was shown to direct the expression of catalytically active heparanase in a mammalian cell expression system. The expressed heparanase was detectable by anti heparanase antibodies in Western blot analysis.

A panel of monochromosomal human/CHO and human/mouse somatic cell hybrids was used to localize the human heparanase gene to human chromosome 4. The newly isolated heparanase sequence can therefore be used to identify a chromosome region harboring a human heparanase gene in a chromosome spread.

The hpa cDNA was then used as a probe to screen a a human genomic library. Several phages were positive. These phages were analyzed and were found to cover most of the hpa locus, except for a small portion which was recovered by bridging PCR. The hpa locus covers about

RT-PCR performed on a variety of cells revealed alternatively spliced *hpa* transcripts.

The amino acid sequence of human heparanase was used to search for homologous sequences in the DNA and protein databases. Several human EST's were identified, as well as mouse sequences highly homologous to human heparanase. The following mouse EST's were identified AA177901, AA674378, AA67997, AA047943, AA690179, AI122034, all sharing an identical sequence and correspond to amino acids 336-543 of the human heparanase sequence. The entire mouse heparanase cDNA was cloned, based on the nucleotide sequence of the mouse EST's using Marathon cDNA libraries. The mouse and the human *hpa* genes share an average homology of 78 % between the nucleotide sequences and 81 % similarity between the deduced amino acid sequences. *hpa* homologous sequences from rat were also uncovered (EST's AI060284 and AI237828).

Homology search of heparanase amino acid sequence against the DNA and the protein databases and prediction of its protein secondary structure enabled to identify candidate amino acids that participate in the heparanase active site.

Expression of *hpa* antisense in mammalian cell lines resulted in about five fold decrease in the number of recoverable cells as compared to

Human *Hpa* cDNA was shown to hybridize with genomic DNAs of a variety of mammalian species and with an avian.

The human and mouse *hpa* promoters were identified and the human promoter was tested positive in directing the expression of a reporter gene.

Thus, according to the present invention there is provided an isolated nucleic acid comprising a genomic, complementary or composite polynucleotide sequence encoding a polypeptide having heparanase catalytic activity.

The phrase "composite polynucleotide sequence" refers to a sequence which includes exonal sequences required to encode the polypeptide having heparanase activity, as well as any number of intronal sequences. The intronal sequences can be of any source and typically will include conserved splicing signal sequences. Such intronal sequences may further include cis acting expression regulatory elements.

The term "heparanase catalytic activity" or its equivalent term "heparanase activity" both refer to a mammalian endoglycosidase hydrolyzing activity which is specific for heparan or heparan sulfate proteoglycan substrates, as opposed to the activity of bacterial enzymes (heparinase I, II and III) which degrade heparin or heparan sulfate by means of β-elimination (37).

polynucleotide and a polypeptide encoded by it, such as SEQ ID NO:

42, 43 or a portion thereof at 68 °C in 6 x SSC, 1 % SDS, 5 x Denharts, 10 % dextran sulfate, 100 µg ml salmon sperm DNA, and ^{32}p labeled probe and wash at 68 °C with 3, 2, 1, 0.5 or 0.1 x SSC and 0.1 % SDS.

According to another preferred embodiment of the present invention the polynucleotide or a portion thereof is at least 60 %, preferably at least 65 %, more preferably at least 70 %, more preferably at least 75 %, more preferably at least 80 %, more preferably at least 85 %, more preferably at least 90 %, most preferably, 95-100 % identical with SEQ ID NOs: 9, 13, 42, 43 or portions thereof as determined using the Bestfit procedure of the DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the university of Wisconsin (gap creation penalty - 12, gap extension penalty - 4 - which are the default parameters).

According to another preferred embodiment of the present invention the polypeptide encoded by the polynucleotide sequence is as set forth in SEQ ID NOs:10, 14, 44 or portions thereof having heparanase catalytic activity. Such portions are expected to include amino acids Asp-Glu 224-225 (SEQ ID NO:10), which can serve as proton donors and glutamic acid 343 or 396 which can serve as a nucleophile.

According to another preferred embodiment of the present invention the polypeptide encoded by the polynucleotide sequence is at least 60 %,

least [illegible] preferably at least [illegible]

more preferably at least 90 %, most preferably, 95-100 % homologous (both similar and identical acids) to SEQ ID NOs:10, 14, 44 or portions thereof as determined with the Smith-Waterman algorithm, using the Bioaccelerator platform developed by Compugene (gapop: 10.0, gapext: 0.5, matrix: blosum62, see also the description to Figure 17).

Further according to the present invention there is provided a nucleic acid construct comprising the isolated nucleic acid described herein. The construct may and preferably further include an origin of replication and trans regulatory elements, such as promoter and enhancer sequences.

The construct or vector can be of any type. It may be a phage which infects bacteria or a virus which infects eukaryotic cells. It may also be a plasmid, phagemid, cosmid, baemid or an artificial chromosome.

Further according to the present invention there is provided a host cell comprising the nucleic acid construct described herein. The host cell can be of any type. It may be a prokaryotic cell, an eukaryotic cell, a cell line, or a cell as a portion of an organism. The polynucleotide encoding heparanase can be permanently or transiently present in the cell. In other words, genetically modified cells obtained following stable or transient transfection, transformation or transduction are all within the scope of the present invention. The polynucleotide can be present in the cell in low copy

integrated in one or more chromosomes at any location or be present as an extrachromosomal material.

The present invention is further directed at providing a heparanase over-expression system which includes a cell overexpressing heparanase catalytic activity. The cell may be a genetically modified host cell transiently or stably transfected or transformed with any suitable vector which includes a polynucleotide sequence encoding a polypeptide having heparanase activity and a suitable promoter and enhancer sequences to direct over-expression of heparanase. However, the overexpressing cell may also be a product of an insertion (e.g., via homologous recombination) of a promoter and/or enhancer sequence downstream to the endogenous heparanase gene of the expressing cell, which will direct over-expression from the endogenous gene.

The term "over-expression" as used herein in the specification and claims below refers to a level of expression which is higher than a basal level of expression typically characterizing a given cell under otherwise identical conditions.

According to another aspect the present invention provides an antisense oligonucleotide comprising a polynucleotide or a polynucleotide analog of at least 10, preferably 11-15, more preferably 16-17, more

preferably [illegible] cell, hybridizable [illegible] coded by nucleic

conditions, with a portion of a polynucleotide strand encoding a polypeptide having heparanase catalytic activity. The antisense oligonucleotide can be used for downregulating heparanase activity by *in vivo* administration thereof to a patient. As such, the antisense oligonucleotide according to the present invention can be used to treat types of cancers which are characterized by impaired (over) expression of heparanase, and are dependent on the expression of heparanase for proliferating or forming metastases.

The antisense oligonucleotide can be DNA or RNA or even include nucleotide analogs, examples of which are provided in the Background section hereinabove. The antisense oligonucleotide according to the present invention can be synthetic and is preferably prepared by solid phase synthesis. In addition, it can be of any desired length which still provides specific base pairing (e.g., 8 or 10, preferably more, nucleotides long) and it can include mismatches that do not hamper base pairing under physiological conditions.

Further according to the present invention there is provided a pharmaceutical composition comprising the antisense oligonucleotide herein described and a pharmaceutically acceptable carrier. The carrier can be, for example, a liposome loadable with the antisense oligonucleotide.

[illegible] oligonucleotide further include [illegible] sequence. The

ribozyme sequence serves to cleave a heparanase RNA molecule to which the antisense oligonucleotide binds, to thereby downregulate heparanase expression.

Further according to the present invention there is provided an antisense nucleic acid construct comprising a promoter sequence and a polynucleotide sequence directing the synthesis of an antisense RNA sequence of at least 10 bases being hybridizable *in vivo*, under physiological conditions, with a portion of a polynucleotide strand encoding a polypeptide having heparanase catalytic activity. Like the antisense oligonucleotide, the antisense construct can be used for downregulating heparanase activity by *in vivo* administration thereof to a patient. As such, the antisense construct, like the antisense oligonucleotide, according to the present invention can be used to treat types of cancers which are characterized by impaired (over) expression of heparanase, and are dependent on the expression of heparanase for proliferating or forming metastases.

Thus, further according to the present invention there is provided a pharmaceutical composition comprising the antisense construct herein described and a pharmaceutically acceptable carrier. The carrier can be, for example, a liposome loadable with the antisense construct.

Formulations for topical administration may include, but are not

powder or oily bases, thickeners and the like may be necessary or desirable. Coated condoms, stents, active pads, and other medical devices may also be useful. Compositions for oral administration include powders or granules, suspensions or solutions in water or non-aqueous media, sachets, capsules or tablets. Thickeners, diluents, flavorings, dispersing aids, emulsifiers or binders may be desirable. Formulations for parenteral administration may include, but are not limited to, sterile aqueous solutions which may also contain buffers, diluents and other suitable additives.

Dosing is dependent on severity and responsiveness of the condition to be treated, but will normally be one or more doses per day, week or month with course of treatment lasting from several days to several months or until a cure is effected or a diminution of disease state is achieved. Persons ordinarily skilled in the art can easily determine optimum dosages, dosing methodologies and repetition rates.

Further according to the present invention there is provided a nucleic acid construct comprising a polynucleotide sequence functioning as a promoter, the polynucleotide sequence is derived from SEQ ID NO:42 and includes at least nucleotides 2135-2635, preferably 2235-2635, more preferably 2335-2635, more preferably 2435-2635, most preferably 2535-2635 thereof, or SEQ ID NO:43 and includes at least nucleotides

follows to direct the synthesis of a reporter gene in transformed cells. Thus, further according to the present invention there is provided a method of expressing a polynucleotide sequence comprising the step of ligating the polynucleotide sequence downstream to either of the promoter sequences described herein. Heparanase promoters can be isolated from a variety of mammalian an other species by cloning genomic regions present 5' to the coding sequence thereof. This can be readily achievable by one ordinarily skilled in the art using the heparanase polynucleotides described herein, which are shown in the Examples section that follows to participate in efficient cross species hybridization.

Further according to the present invention there is provided a recombinant protein comprising a polypeptide having heparanase catalytic activity. The protein according to the present invention include modifications known as post translational modifications, including, but not limited to, proteolysis (e.g., removal of a signal peptide and of a pro- or preprotein sequence), methionine modification, glycosylation, alkylation (e.g., methylation), acetylation, etc. According to preferred embodiments the polypeptide includes at least a portion of SEQ ID NOs:10, 14 or 44, the portion has heparanase catalytic activity. According to preferred embodiments of the present invention the protein is encoded by any of the

invention there is provided a pharmaceutical composition comprising, as an active ingredient, the recombinant protein described herein.

The recombinant protein may be purified by any conventional protein purification procedure close to homogeneity and/or be mixed with additives. The recombinant protein may be manufactured using any of the genetically modified cells described above, which include any of the expression nucleic acid constructs described herein. The recombinant protein may be in any form. It may be in a crystallized form, a dehydrated powder form or in solution. The recombinant protein may be useful in obtaining pure heparanase, which in turn may be useful in eliciting anti-heparanase antibodies, either poly or monoclonal antibodies, and as a screening active ingredient in an anti-heparanase inhibitors or drugs screening assay or system.

Further according to the present invention there is provided a method of identifying a chromosome region harboring a human heparanase gene in a chromosome spread. the method is executed implementing the following method steps, in which in a first step the chromosome spread (either interphase or metaphase spread) is hybridized with a tagged polynucleotide probe encoding heparanase. The tag is preferably a fluorescent tag. In a second step according to the method the chromosome spread is washed, associated with the hybridized tagged polynucleotide probe are scattered for,

wherein detected signals being indicative of a chromosome region harboring the human heparanase gene. One ordinarily skilled in the art would know how to use the sequences disclosed herein in suitable labeling reactions and how to use the tagged probes to detect, using *in situ* hybridization, a chromosome region harboring a human heparanase gene.

Further according to the present invention there is provided a method of *in vivo* eliciting anti-heparanase antibodies comprising the steps of administering a nucleic acid construct including a polynucleotide segment corresponding to at least a portion of SEQ ID NOs:9, 13 or 43 and a promoter for directing the expression of said polynucleotide segment *in vivo*. Accordingly, there is provided also a DNA vaccine for *in vivo* eliciting anti-heparanase antibodies comprising a nucleic acid construct including a polynucleotide segment corresponding to at least a portion of SEQ ID NOs:9, 13 or 43 and a promoter for directing the expression of said polynucleotide segment *in vivo*. The vaccine optionally further includes a pharmaceutically acceptable carrier, such as a virus, liposome or an antigen presenting cell. Alternatively, the vaccine is employed as a naked DNA vaccine

The present invention can be used to develop treatments for various diseases, to develop diagnostic assays for these diseases and to provide new

Specifically, the present invention can be used to develop new drugs to inhibit tumor cell metastasis, inflammation and autoimmunity. The identification of the *hpa* gene encoding for the heparanase enzyme enables the production of a recombinant enzyme in heterologous expression systems.

Furthermore, the present invention can be used to modulate bioavailability of heparin-binding growth factors, cellular responses to heparin-binding growth factors (e.g., bFGF, VEGF) and cytokines (e.g., IL-8), cell interaction with plasma lipoproteins, cellular susceptibility to viral, protozoa and some bacterial infections, and disintegration of neurodegenerative plaques. Recombinant heparanase offers a potential treatment for wound healing, angiogenesis, restenosis, atherosclerosis, inflammation, neurodegenerative diseases (such as, for example, Genstmann-Straussler Syndrome, Creutzfeldt-Jakob disease, Scrape and Alzheimer's disease) and certain viral and some bacterial and protozoa infections. Recombinant heparanase can be used to neutralize plasma heparin, as a potential replacement of protamine.

As used herein, the term "modulate" includes substantially inhibiting, slowing or reversing the progression of a disease, substantially ameliorating clinical symptoms of a disease or condition, or substantially preventing the

therefore include an agent which may modulate a disease or condition

Modulation of viral, protozoa and bacterial infections includes any effect which substantially interrupts, prevents or reduces any viral, bacterial or protozoa activity and/or stage of the virus, bacterium or protozoon life cycle, or which reduces or prevents infection by the virus, bacterium or protozoon in a subject, such as a human or lower animal.

As used herein, the term "wound" includes any injury to any portion of the body of a subject including, but not limited to, acute conditions such as thermal burns, chemical burns, radiation burns, burns caused by excess exposure to ultraviolet radiation such as sunburn, damage to bodily tissues such as the perineum as a result of labor and childbirth, including injuries sustained during medical procedures such as episiotomies, trauma-induced injuries including cuts, those injuries sustained in automobile and other mechanical accidents, and those caused by bullets, knives and other weapons, and post-surgical injuries, as well as chronic conditions such as pressure sores, bedsores, conditions related to diabetes and poor circulation, and all types of acne, etc.

Anti-heparanase antibodies, raised against the recombinant enzyme, would be useful for immunodetection and diagnosis of micrometastases, autoimmune lesions and renal failure in biopsy specimens, plasma samples, and body fluids. Such antibodies may also serve as neutralizing agents for

The genomic heparanase sequences described herein can be used to construct knock-in and knock-out constructs. Such constructs include a fragment of 10-20 Kb of a heparanase locus and a negative and a positive selection markers and can be used to provide heparanase knock-in and knock-out animal models by methods known to the skilled artisan. Such animal models can be used for studying the function of heparanase in developmental processes, and in normal as well as pathological processes. They can also serve as an experimental model for testing drugs and gene therapy protocols. The complementary heparanase sequence (cDNA) can be used to derive transgenic animals, overexpressing heparanase for same. Alternatively , if cloned in the antisense orientation, the complementary heparanase sequence (cDNA) can be used to derive transgenic animals under-expressing heparanase for same.

The heparanase promoter sequences described herein and other cis regulatory elements linked to the heparanase locus can be used to regulated the expression of genes. For example, these promoters can be used to direct the expression of a cytotoxic protein, such as TNF, in tumor cells. It will be appreciated that heparanase itself is abnormally expressed under the control of its own promoter and other cis acting elements in a variety of tumors, and its expression is correlated with metastasis. It is also

heparanase cDNA can be used for the same purpose, as it is known that

introns, especially upstream introns include cis acting element which affect expression. A heparanase promoter fused to a reporter protein can be used to study/monitor its activity.

The polynucleotide sequences described herein can also be used to provide DNA vaccines which will elicit in vivo anti heparanase antibodies. Such vaccines can therefore be used to combat inflammatory and cancer.

Antisense oligonucleotides derived according to the heparanase sequences described herein, especially such oligonucleotides supplemented with ribozyme activity, can be used to modulate heparanase expression. Such oligonucleotides can be from the coding region, from the introns or promoter specific. Antisense heparanase nucleic acid constructs can similarly function, as well known in the art.

The heparanase sequences described herein can be used to study the catalytic mechanism of heparanase. Carefully selected site directed mutagenesis can be employed to provide modified heparanase proteins having modified characteristics in terms of, for example, substrate specificity, sensitivity to inhibitors, etc.

While studying heparanase expression in a variety of cell types alternatively spliced transcripts were identified. Such transcripts if found characteristic of certain pathological conditions can be used as markers for

Additional objects, advantages, and novel features of the present invention will become apparent to one ordinarily skilled in the art upon examination of the following examples, which are not intended to be limiting. Additionally, each of the various embodiments and aspects of the present invention as delineated hereinabove and as claimed in the claims section below finds experimental support in the following examples.

EXAMPLES

Generally, the nomenclature used herein and the laboratory procedures in recombinant DNA technology described below are those well known and commonly employed in the art. Standard techniques are used for cloning, DNA and RNA isolation, amplification and purification. Generally enzymatic reactions involving DNA ligase, DNA polymerase, restriction endonucleases and the like are performed according to the manufacturers' specifications. These techniques and various other techniques are generally performed according to Sambrook et al., Molecular Cloning--A Laboratory Manual, Cold Spring Harbor Laboratory, Cold Spring Harbor, N.Y. (1989), which is incorporated herein by reference. Other general references are provided throughout this document. The procedures therein are believed to be well known in the art and are provided for the convenience of the reader.

The following protocols and experimental details are referenced in the Examples that follow:

Purification and characterization of heparanase from a human hepatoma cell line and human placenta: A human hepatoma cell line (Sk-hep-1) was chosen as a source for purification of a human tumor-derived heparanase. Purification was essentially as described in U.S. Pat. No. 5,362,641 to Fuks, which is incorporated by reference as if fully set forth herein. Briefly, 500 liter, $5x10^{11}$ cells were grown in suspension and the heparanase enzyme was purified about 240,000 fold by applying the following steps: (i) cation exchange (CM-Sephadex) chromatography performed at pH 6.0, 0.3-1.4 M NaCl gradient; (ii) cation exchange (CM-Sephadex) chromatography performed at pH 7.4 in the presence of 0.1% CHAPS, 0.3-1.1 M NaCl gradient; (iii) heparin-Sepharose chromatography performed at pH 7.4 in the presence of 0.1% CHAPS, 0.35-1.1 M NaCl gradient; (iv) ConA-Sepharose chromatography performed at pH 6.0 in buffer containing 0.1 % CHAPS and 1 M NaCl, elution with 0.25 M α-methyl mannoside; and (v) HPLC cation exchange (Mono-S) chromatography performed at pH 7.4 in the presence of 0.1 % CHAPS, 0.25-1 M NaCl gradient.

precipitate subjected to SDS polyacrylamide gel electrophoresis and of

tryptic digestion and reverse phase HPLC. Tryptic peptides of the purified protein were separated by reverse phase HPLC (C8 column) and homogeneous peaks were subjected to amino acid sequence analysis.

The purified enzyme was applied to reverse phase HPLC and subjected to N-terminal amino acid sequencing using the amino acid sequencer (Applied Biosystems).

Cells: Cultures of bovine corneal endothelial cells (BCECs) were established from steer eyes as previously described (19, 38). Stock cultures were maintained in DMEM (1 g glucose/liter) supplemented with 10 % newborn calf serum and 5 % FCS. bFGF (1 ng/ml) was added every other day during the phase of active cell growth (13, 14).

Preparation of dishes coated with ECM: BCECs (second to fifth passage) were plated into 4-well plates at an initial density of 2 x 10^5 cells/ml, and cultured in sulfate-free Fisher medium plus 5 % dextran T-40 for 12 days. $Na_2{}^{35}SO_4$ (25 µCi/ml) was added on day 1 and 5 after seeding and the cultures were incubated with the label without medium change. The subendothelial ECM was exposed by dissolving (5 min., room temperature) the cell layer with PBS containing 0.5 % Triton X-100 and 20 mM NH_4OH, followed by four washes with PBS. The ECM remained intact, free of cellular debris and firmly attached to the entire area of the tissue culture

To prepare soluble sulfate labeled proteoglycans (peak I material), the ECM was digested with trypsin (25 μg/ml, 6 h, 37 °C), the digest was concentrated by reverse dialysis and the concentrated material was applied onto a Sepharose 6B gel filtration column. The resulting high molecular weight material (K_{av} < 0.2, peak I) was collected. More than 80 % of the labeled material was shown to be composed of heparan sulfate proteoglycans (11, 39).

Heparanase activity: Cells (1 x 10^6/35-mm dish), cell lysates or conditioned media were incubated on top of ^{35}S-labeled ECM (18 h, 37 °C) in the presence of 20 mM phosphate buffer (pH 6.2). Cell lysates and conditioned media were also incubated with sulfate labeled peak I material (10-20 μl). The incubation medium was collected, centrifuged (18,000 x g, 4 °C, 3 min.), and sulfate labeled material analyzed by gel filtration on a Sepharose CL-6B column (0.9 x 30 cm). Fractions (0.2 ml) were eluted with PBS at a flow rate of 5 ml/h and counted for radioactivity using Bio-fluor scintillation fluid. The excluded volume (V_0) was marked by blue dextran and the total included volume (V_t) by phenol red. The latter was shown to comigrate with free sulfate (7, 11, 23). Degradation fragments of HS side chains were eluted from Sepharose 6B at 0.5 < K_{av} < 0.8 (peak II) (7, 11, 23). A nearly intact HSPG released from ECM by trypsin - and,

ranged from 85 to 95 % in different experiments (11). Each experiment was performed at least three times and the variation of elution positions (Kav values) did not exceed +/- 15 %.

Cloning of hpa cDNA: cDNA clones 257548 and 260138 were obtained from the I.M.A.G.E Consortium (2130 Memorial Parkway SW, Hunstville, AL 35801). The cDNAs were originally cloned in *Eco*RI and *Not*I cloning sites in the plasmid vector pT3T7D-Pac. Although these clones are reported to be somewhat different, DNA sequencing demonstrated that these clones are identical to one another. Marathon RACE (rapid amplification of cDNA ends) human placenta (poly-A) cDNA composite was a gift of Prof. Yossi Shiloh of Tel Aviv University. This composite is vector free, as it includes reverse transcribed cDNA fragments to which double, partially single stranded adapters are attached on both sides. The construction of the specific composite employed is described in reference 39a.

Amplification of hp3 PCR fragment was performed according to the protocol provided by Clontech laboratories. The template used for amplification was a sample taken from the above composite. The primers used for amplification were:

First step: 5'-primer: AP1: 5'-CCATCCTAATACGACTCACT

Second step: nested 5'-primer: AP2: 5'-ACTCACTATAGGGCTCG AGCGGC-3', SEQ ID NO:3; nested 3'- primer: HPL171: 5'-GCATCTTAGCCGTCTTTCTTCG-3', SEQ ID NO:4. The HPL229 and HPL171 were selected according to the sequence of the EST clones. They include nucleotides 933-956 and 876-897 of SEQ ID NO:9, respectively.

PCR program was 94 °C - 4 min., followed by 30 cycles of 94 °C - 40 sec., 62 °C - 1 min., 72 °C - 2.5 min. Amplification was performed with Expand High Fidelity (Boehringer Mannheim). The resulting ca. 900 bp hp3 PCR product was digested with *Bfr*I and *Pvu*II. Clone 257548 (p*hpa*1) was digested with *Eco*RI, followed by end filling and was then further digested with *Bfr*I. Thereafter the *Pvu*II - *Bfr*I fragment of the hp3 PCR product was cloned into the blunt end - *Bfr*I end of clone p*hpa*1 which resulted in having the entire cDNA cloned in pT3T7-pac vector, designated p*hpa*2.

RT-PCR: RNA was prepared using TRI-Reagent (Molecular research center Inc.) according to the manufacturer instructions. 1.25 µg were taken for reverse transcription reaction using MuMLV Reverse transcriptase (Gibco BRL) and Oligo $(dT)_{15}$ primer, SEQ ID NO:5, (Promega). Amplification of the resultant first strand cDNA was performed with *Taq* polymerase (Promega). The following primers were

HPL229: 5'-TTCGAATCCCAAGAAGGAAATCAAG-3', SEQ ID NO:6;

nucleotides 372-394 in SEQ ID NOs:9 or 11.

HPL-229: 5'-GTAGTGATGCCATGTAACTGAATC-3', SEQ ID NO:7, nucleotides 933-956 in SEQ ID NOs:9 or 11.

PCR program: 94 °C - 4 min., followed by 30 cycles of 94 °C - 40 sec., 62 °C - 1 min., 72 °C - 1 min.

Alternatively, total RNA was prepared from cell cultures using Tri-reagent (Molecular Research Center, Inc.) according to the manufacturer recommendation. Poly A+ RNA was isolated from total RNA using mRNA separator (Clontech). Reverse transcription was performed with total RNA using Superscript II (GibcoBRL). PCR was performed with Expand high fidelity (Boehringer Mannheim). Primers used for amplification were as follows:

Hpu-685, 5'-GAGCAGCCAGGTGAGCCCAAGAT-3', SEQ ID NO:24

Hpu-355, 5'-TTCGATCCCAAGAAGGAATCAAC-3', SEQ ID NO:25

Hpu 565, 5'-AGCTCTGTAGATGTGCTATACAC-3', SEQ ID NO:26

Hpl 967, 5'-TCAGATGCAAGCAGCAACTTTGGC-3', SEQ ID NO:27

Hpl 171, 5'-GCATCTTAGCCGTCTTTCTTCG-3', SEQ ID NO:28

Hpl 229, 5'-GTAGTGATGCCATGTAACTGAATC-3', SEQ ID NO:29

PCR reaction was performed as follows: 94 °C 3 minutes, followed by 32 cycles of 94 °C 40 seconds, 64 °C 1 minute, 72 °C 3 minutes, and one

Expression of recombinant heparanase in insect cells: Cells, High Five and Sf21 insect cell lines were maintained as monolayer cultures in SF900II-SFM medium (GibcoBRL).

Recombinant Baculovirus: Recombinant virus containing the *hpa* gene was constructed using the Bac to Bac system (GibcoBRL). The transfer vector pFastBac was digested with *Sal*I and *Not*I and ligated with a 1.7 kb fragment of p*hpa*2 digested with *Xho*I and *Not*I. The resulting plasmid was designated pFast*hpa*2. An identical plasmid designated pFast*hpa*4 was prepared as a duplicate and both independently served for further experimentations. Recombinant bacmid was generated according to the instructions of the manufacturer with pFast*hpa*2, pFast*hpa*4 and with pFastBac. The latter served as a negative control. Recombinant bacmid DNAs were transfected into Sf21 insect cells. Five days after transfection recombinant viruses were harvested and used to infect High Five insect cells, 3 x 10^6 cells in T-25 flasks. Cells were harvested 2 - 3 days after infection. 4 x 10^6 cells were centrifuged and resuspended in a reaction buffer containing 20 mM phosphate citrate buffer, 50 mM NaCl. Cells underwent three cycles of freeze and thaw and lysates were stored at -80 °C. Conditioned medium was stored at 4 °C.

Partial purification of recombinant heparanase: Partial

heparin-Sepharose column chromatography followed by Superdex

column gel filtration. Culture medium (150 ml) of Sf21 cells infected with pFhpa4 virus was subjected to heparin-Sepharose chromatography. Elution of 1 ml fractions was performed with 0.35 - 2 M NaCl gradient in presence of 0.1 % CHAPS and 1 mM DTT in 10 mM sodium acetate buffer, pH 5.0. A 25 µl sample of each fraction was tested for heparanase activity. Heparanase activity was eluted at the range of 0.65 - 1.1 M NaCl (fractions 18-26, Figure 10a). 5 µl of each fraction was subjected to 15 % SDS-polyacrylamide gel electrophoresis followed by silver nitrate staining. Active fractions eluted from heparin-Sepharose (Figure 10a) were pooled and concentrated (x 6) on YM3 cut-off membrane. 0.5 ml of the concentrated material was applied onto 30 ml Superdex 75 FPLC column equilibrated with 10 mM sodium acetate buffer, pH 5.0, containing 0.8 M NaCl, 1 mM DTT and 0.1 % CHAPS. Fractions (0.56 ml) were collected at a flow rate of 0.75 ml/min. Aliquots of each fraction were tested for heparanase activity and were subjected to SDS-polyacrylamide gel electrophoresis followed by silver nitrate staining (Figure 11b).

PCR amplification of genomic DNA: 94 °C 3 minutes, followed by 32 cycles of 94 °C 45 seconds, 64 °C 1 minute, 68 °C 5 minutes, and one cycle at 72 °C, 7 minutes. Primers used for amplification of genomic DNA included:

GHpI-Lo [illegible] (SEQ ID NO:51)

Screening of genomic libraries: A human genomic library in Lambda phage EMBLE3 SP6/T7 (Clontech, Paulo Alto, CA) was screened. 5×10^5 plaques were plated at 5×10^4 pfu/plate on NZCYM agar/top agarose plates. Phages were absorbed on nylon membranes in duplicates (Qiagen). Hybridization was performed at 65 °C in 5 x SSC, 5 x Denhart's, 10 % dextran sulfate, 100 µg/ml Salmon sperm, ^{32}p labeled probe (10^6 cpm/ml). A 1.6 kb fragment, containing the entire *hpa* cDNA was labeled by random priming (Boehringer Mannheim). Following hybridization membranes were washed once with 2 x SSC, 0.1 % SDS at 65 °C for 20 minutes, and twice with 0.2 x SSC, 0.1 % SDS at 65 °C for 15 minutes. Hybridizing plaques were picked, and plated at 100 pfu/plate. Hybridization was performed as above and single isolated positive plaques were picked.

Phage DNA was extracted using a Lambda DNA extraction kit (Qiagen). DNA was digested with *Xho*I and *Eco*RI, separated on 0.7 % agarose gel and transferred to nylon membrane Hybond N+ (Amersham). Hybridization and washes were performed as above.

cDNA Sequence analysis: Sequence determinations were performed with vector specific and gene specific primers, using an automated DNA sequencer (Applied Biosystems, model 373A). Each nucleotide was read

Genomic sequence analysis: Large-scale sequencing was performed by Commonwealth Biotechnology Incorporation.

Isolation of mouse hpa: Mouse *hpa* cDNA was amplified from either Marathon ready cDNA library of mouse embryo or from mRNA isolated from mouse melanoma cell line BL6, using the Marathon RACE kit from Clontech. Both procedures were performed according to the manufacturer's recommendation.

Primers used for PCR amplification of mouse hpa:

Mhpl773 5'-CCACACTGAATGTAATACTGAAGTG-3', SEQ ID NO:32

MHpl736 5'-CGAAGCTCTGGAACTCGGCAAG-3', SEQ ID NO:33

MHpl83 5'-GCCAGCTGCAAAGGTGTTGGAC-3', SEQ ID NO:34

Mhpl152 5'-AACACCTGCCTCATCACGACTTC-3', SEQ ID NO:35

Mhpl114 5'-GCCAGGCTGGCGTCGATGGTGA-3', SEQ ID NO:36

MHpl103 5'-GTCGATGGTGATGGACAGGAAC-3', SEQ ID NO:37

Ap1 5'-GTAATACGACTCACTATAGGGC-3', SEQ ID NO:38 - (Genome walker)

Ap2 5'-ACTATAGGGCACGCGTGGT-3', SEQ ID NO:39 - (Genome walker)

Ap1 5'-CCATCCTAATACGACTCACTATAGGGC-3', SEQ ID NO:40 - (Marathon RACE)

(Marathon RACE)

Southern analysis of genomic DNA: Genomic DNA was extracted from animal or from human blood using Blood and cell culture DNA maxi kit (Qiagene). DNA was digested with *Eco*RI, separated by gel electrophoresis and transferred to a nylon membrane Hybond N+ (Amersham). Hybridization was performed at 68 °C in 6 x SSC, 1 % SDS, 5 x Denharts, 10 % dextran sulfate, 100 µg/ml salmon sperm DNA, and ^{32}p labeled probe. A 1.6 kb fragment, containing the entire *hpa* cDNA was used as a probe. Following hybridization, the membrane was washed with 3 x SSC, 0.1 % SDS, at 68 °C and exposed to X-ray film for 3 days. Membranes were then washed with 1 x SSC, 0.1 % SDS, at 68 °C and were reexposed for 5 days.

Construction of hpa promoter-GFP expression vector: Lambda DNA of phage L3, was digested with *Sac*I and *Bgl*II, resulting in a 1712 bp fragment which contained the *hpa* promoter (877-2688 of SEQ ID NO:42). The pEGFP-1 plasmid (Clontech) was digested with *Bgl*II and *Sac*I and ligated with the 1712 bp fragment of the *hpa* promoter sequence. The resulting plasmid was designated phpEGL. A second *hpa* promoter-GFP plasmid was constructed containing a shorter fragment of the *hpa* promoter region: phpEGL was digested with *Hind*III, and the resulting 1095 bp fragment (nucleotides 1593-2688 of SEQ ID NO:42) was ligated with

Computer analysis of sequences: Homology searches were performed using several computer servers, and various databases. Blast 2.0 service, at the NCBI server was used to screen the protein database swplus and DNA databases such as GenBank, EMBL, and the EST databases. Blast 2.0 search was performed using the basic search option of the NCBI server. Sequence analysis and alignments were done using the DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the university of Wisconsin. Alignments of two sequences were performed using Bestfit (gap creation penalty - 12, gap extension penalty - 4). Protein homology search was performed with the Smith-Waterman algorithm, using the Bioaccelerator platform developed by Compugene. The protein database swplus was searched using the following parameters: gapop: 10.0, gapext: 0.5, matrix: blosum62. Blocks homology was performed using the Blocks WWW server developed at Fred Hutchinson Cancer Research Center in Seattle, Washington, USA. Secondary structure prediction was performed using the PHD server Profile network Prediction Heidelberg. Fold recognition (threading) was performed using the UCLA-DOE structure prediction server. The method used for prediction was gonnet+predss. Alignment of three sequences was performed using the pileup application (gap creation penalty - 5, gap

TSSG programs (BCM Search Launcher Human Genome Center, Baylor College of Medicine, Houston TX).

EXAMPLE 1

Cloning of human hpa cDNA

Purified fraction of heparanase isolated from human hepatoma cells (SK-hep-1) was subjected to tryptic digestion and microsequencing. EST (Expressed Sequence Tag) databases were screened for homology to the back translated DNA sequences corresponding to the obtained peptides. Two EST sequences (accession Nos. N41349 and N45367) contained a DNA sequence encoding the peptide YGPDVGQPR (SEQ ID NO:8). These two sequences were derived from clones 257548 and 260138 (I.M.A.G.E Consortium) prepared from 8 to 9 weeks placenta cDNA library (Soares). Both clones which were found to be identical contained an insert of 1020 bp which included an open reading frame (ORF) of 973 bp followed by a 3' untranslated region of 27 bp and a Poly A tail. No translation start site (AUG) was identified at the 5' end of these clones.

Cloning of the missing 5' end was performed by PCR amplification of DNA from a placenta Marathon RACE cDNA composite. A 900 bp fragment (designated hp3), partially overlapping with the identified 3'

The joined cDNA fragment, 1721 bp long (SEQ ID NO:9), contained an open reading frame which encodes, as shown in Figure 1 and SEQ ID NO:11, a polypeptide of 543 amino acids (SEQ ID NO:10) with a calculated molecular weight of 61,192 daltons. The 3' end of the partial cDNA inserts contained in clones 257548 and 260138 started at nucleotide G^{721} of SEQ ID NO:9 and Figure 1.

As further shown in Figure 1, there was a single sequence discrepancy between the EST clones and the PCR amplified sequence, which led to an amino acid substitution from Tyr^{246} in the EST to Phe^{246} in the amplified cDNA. The nucleotide sequence of the PCR amplified cDNA fragment was verified from two independent amplification products. The new gene was designated *hpa*.

As stated above, the 3' end of the partial cDNA inserts contained in EST clones 257548 and 260138 started at nucleotide 721 of *hpa* (SEQ ID NO:9). The ability of the *hpa* cDNA to form stable secondary structures, such as stem and loop structures involving nucleotide stretches in the vicinity of position 721 was investigated using computer modeling. It was found that stable stem and loop structures are likely to be formed involving nucleotides 698-724 (SEQ ID NO:9). In addition, a high GC content, up to 70 %, characterizes the 5' end region of the *hpa* gene, as compared to about

To examine the ability of the *hpa* gene product to catalyze degradation of heparan sulfate in an *in vitro* assay the entire open reading frame was expressed in insect cells, using the Baculovirus expression system. Extracts of cells, infected with virus containing the *hpa* gene, demonstrated a high level of heparan sulfate degradation activity, while cells infected with a similar construct containing no *hpa* gene had no such activity, nor did non-infected cells. These results are further demonstrated in the following Examples.

EXAMPLE 2

Degradation of soluble ECM-derived HSPG

Monolayer cultures of High Five cells were infected (72 h, 28 °C) with recombinant Bacoluvirus containing the pFast*hpa* plasmid or with control virus containing an insert free plasmid. The cells were harvested and lysed in heparanase reaction buffer by three cycles of freezing and thawing. The cell lysates were then incubated (18 h, 37 °C) with sulfate labeled, ECM-derived HSPG (peak 1), followed by gel filtration analysis (Sepharose 6B) of the reaction mixture.

As shown in Figure 2, the substrate alone included almost entirely high molecular weight (Mr) material eluted next to V_0 (peak I, fractions

substrate was incubated with lysate of cells that were infected with cells

virus. In contrast, incubation of the HSPG substrate with lysates of cells infected with the *hpa* containing virus resulted in a complete conversion of the high Mr substrate into low Mr labeled degradation fragments (peak II, fractions 22-35, $0.5 < Kav < 0.75$).

Fragments eluted in peak II were shown to be degradation products of heparan sulfate, as they were (i) 5- to 6-fold smaller than intact heparan sulfate side chains (Kav approx. 0.33) released from ECM by treatment with either alkaline borohydride or papain; and (ii) resistant to further digestion with papain or chondroitinase ABC, and susceptible to deamination by nitrous acid (6, 11). Similar results (not shown) were obtained with Sf21 cells. Again, heparanase activity was detected in cells infected with the *hpa* containing virus (pF*hpa*), but not with control virus (pF). This result was obtained with two independently generated recombinant viruses. Lysates of control not infected High Five cells failed to degrade the HSPG substrate.

In subsequent experiments, the labeled HSPG substrate was incubated with medium conditioned by infected High Five or Sf21 cells.

As shown in Figures 3a-b, heparanase activity, reflected by the conversion of the high Mr peak I substrate into the low Mr peak II which represents HS degradation fragments, was found in the culture medium of

pF1 or pF2 viruses. No heparanase activity was detected in the culture medium of control non-infected High Five or Sf21 cells.

The medium of cells infected with the pF*hpa*4 virus was passed through a 50 kDa cut off membrane to obtain a crude estimation of the molecular weight of the recombinant heparanase enzyme. As demonstrated in Figure 4, all the enzymatic activity was retained in the upper compartment and there was no activity in the flow through (<50 kDa) material. This result is consistent with the expected molecular weight of the *hpa* gene product.

In order to further characterize the *hpa* product the inhibitory effect of heparin, a potent inhibitor of heparanase mediated HS degradation (40) was examined.

As demonstrated in Figures 5a-b, conversion of the peak I substrate into peak II HS degradation fragments was completely abolished in the presence of heparin.

Altogether, these results indicate that the heparanase enzyme is expressed in an active form by insect cells infected with Baculovirus containing the newly identified human *hpa* gene.

EXAMPLE 3

Degradation of HSPG in intact ECM

Next, the ability of intact infected insect cells to degrade HS in intact, naturally produced ECM was investigated. For this purpose, High Five or Sf21 cells were seeded on metabolically sulfate labeled ECM followed by infection (48 h, 28 °C) with either the pF*hpa*4 or control pF2 viruses. The pH of the medium was then adjusted to pH 6.2-6.4 and the cells further incubated with the labeled ECM for another 48 h at 28 °C or 24 h at 37 °C. Sulfate labeled material released into the incubation medium was analyzed by gel filtration on Sepharose 6B.

As shown in Figures 6a-b and 7a-b, incubation of the ECM with cells infected with the control pF2 virus resulted in a constant release of labeled material that consisted almost entirely (>90%) of high Mr fragments (peak I) eluted with or next to V_O. It was previously shown that a proteolytic activity residing in the ECM itself and/or expressed by cells is responsible for release of the high Mr material (6). This nearly intact HSPG provides a soluble substrate for subsequent degradation by heparanase, as also indicated by the relatively large amount of peak I material accumulating when the heparanase enzyme is inhibited by heparin (6, 7, 12, Figure 9). On the other hand, incubation of the labeled ECM with cells infected with the

[illegible] of the control virus, [illegible] sulfate-labeled fragments (peak II, [illegible]

<Kav< 0.75), regardless of whether the infected cells were incubated with the ECM at 28 °C or 37 °C. Control intact non-infected Sf21 or High Five cells failed to degrade the ECM HS side chains.

In subsequent experiments, as demonstrated in Figures 8a-b, High Five and Sf21 cells were infected (96 h, 28 °C) with pF*hpa*4 or control pF1 viruses and the culture medium incubated with sulfate-labeled ECM. Low Mr HS degradation fragments were released from the ECM only upon incubation with medium conditioned by pF*hpa*4 infected cells. As shown in Figure 9, production of these fragments was abolished in the presence of heparin. No heparanase activity was detected in the culture medium of control, non-infected cells. These results indicate that the heparanase enzyme expressed by cells infected with the pF*hpa*4 virus is capable of degrading HS when complexed to other macromolecular constituents (i.e. fibronectin, laminin, collagen) of a naturally produced intact ECM, in a manner similar to that reported for highly metastatic tumor cells or activated cells of the immune system (6, 7).

EXAMPLE 4

Purification of recombinant human heparanase

The recombinant heparanase was partially purified from medium of

[illegible] by ion exchange of the pooled active fractions over an FPLC

Superdex 75 column (Figure 11a). A ~ 63 kDa protein was observed, whose quantity, as was detected by silver stained SDS-polyacrylamide gel electrophoresis, correlated with heparanase activity in the relevant column fractions (Figures 10b and 11b, respectively). This protein was not detected in the culture medium of cells infected with the control pF1 virus and was subjected to a similar fractionation on heparin-Sepharose (not shown).

EXAMPLE 5

Expression of the human hpa cDNA in various cell types, organs and tissues

Referring now to Figures 12a-e, RT-PCR was applied to evaluate the expression of the *hpa* gene by various cell types and tissues. For this purpose, total RNA was reverse transcribed and amplified. The expected 585 bp long cDNA was clearly demonstrated in human kidney, placenta (8 and 11 weeks) and mole tissues, as well as in freshly isolated and short termed (1.5-48 h) cultured human placental cytotrophoblastic cells (Figure 12a), all known to express a high heparanase activity (41). The *hpa* transcript was also expressed by normal human neutrophils (Figure 12b). In contrast, there was no detectable expression of the *hpa* mRNA in embryonic human muscle tissue, thymus, heart and adrenal (Figure 12b). The *hpa* gene

breast carcinoma (435, 231), melanoma and megakaryocytic (DAMI, CHRF) human cell lines (Figures 12d-e).

The above described expression pattern of the *hpa* transcript was determined to be in a very good correlation with heparanase activity levels determined in various tissues and cell types (not shown).

EXAMPLE 6

Isolation of an extended 5' end of hpa cDNA from human SK-hep1 cell line

The 5' end of *hpa* cDNA was isolated from human SK-hep1 cell line by PCR amplification using the Marathon RACE (rapid amplification of cDNA ends) kit (Clontech). Total RNA was prepared from SK-hep1 cells using the TRI-Reagent (Molecular research center Inc.) according to the manufacturer instructions. Poly A+ RNA was isolated using the mRNA separator kit (Clonetech).

The Marahton RACE SK-hep1 cDNA composite was constructed according to the manufacturer recommendations. First round of amplification was performed using an adaptor specific primer AP1: 5'-CCATCCTAATACG ACTCACTATAGGGC-3', SEQ ID NO:1, and a *hpa* specific antisense primer hpl-629:

subjected to a second round of amplification using an adaptor specific nested primer AP2: 5'-ACTCACTATAGGGCTCGAGCGGC-3', SEQ ID NO:3, and a *hpa* specific antisense nested primer hpl-666 5'-AGGCTTCGAGCGCAGCAGCAT-3', SEQ ID NO:18, corresponding to nucleotides 83-63 of SEQ ID NO:9. The PCR program was as follows: a hot start of 94 °C for 1 minute, followed by 30 cycles of 90 °C - 30 seconds, 68 °C - 4 minutes. The resulting 300 bp DNA fragment was extracted from an agarose gel and cloned into the vector pGEM-T Easy (Promega). The resulting recombinant plasmid was designated pHPSK1.

The nucleotide sequence of the pHPSK1 insert was determined and it was found to contain 62 nucleotides of the 5' end of the placenta *hpa* cDNA (SEQ ID NO:9) and additional 178 nucleotides upstream, the first 178 nucleotides of SEQ ID NOs:13 and 15.

A single nucleotide discrepancy was identified between the SK-hep1 cDNA and the placenta cDNA. The "T" derivative at position 9 of the placenta cDNA (SEQ ID NO:9), is replaced by a "C" derivative at the corresponding position 187 of the SK-hep1 cDNA (SEQ ID NO:13).

The discrepancy is likely to be due to a mutation at the 5' end of the placenta cDNA clone as confirmed by sequence analysis of sevsral additional cDNA clones isolated from placenta, which like the SK-hep1

The 5' extended sequence of the SK-hep1 *hpa* cDNA was assembled with the sequence of the *hpa* cDNA isolated from human placenta (SEQ ID NO:9). The assembled sequence contained an open reading frame which encodes, as shown in SEQ ID NOs:14 and 15, a polypeptide of 592 amino acids with a calculated molecular weight of 66,407 daltons. The open reading frame is flanked by 93 bp 5' untranslated region (UTR).

EXAMPLE 7

Isolation of the upstream genomic region of the hpa gene

The upstream region of the *hpa* gene was isolated using the Genome Walker kit (Clontech) according to the manufacturer recommendations. The kit includes five human genomic DNA samples each digested with a different restriction endonuclease creating blunt ends: *Eco*RV, *Sca*I, *Dra*I, *Pvu*II and *Ssp*I.

The blunt ended DNA fragments are ligated to partially single stranded adaptors. The Genomic DNA samples were subjected to PCR amplification using the adaptor specific primer and a gene specific primer. Amplification was performed with Expand High Fidelity (Boehringer Mannheim).

A first round of amplification was performed using the ap1 primer:

5'-AGGCTTCGAGCGCAGCAGCAT-3', SEQ ID NO:18, corresponding to nucleotides 83 - 63 of SEQ ID NO:9. The PCR program was as follows: a hot start of 94 °C - 3 minutes, followed by 36 cycles of 94 °C - 40 seconds, 67 °C - 4 minutes.

The PCR products of the first amplification were diluted 1:50. One µl of the diluted sample was used as a template for a second amplification using a nested adaptor specific primer ap2: 5'-ACTATAGGGCACGCGTGGT-3', SEQ ID NO:20, and a *hpa* specific antisense primer hpl-690, 5'-CTTGGGCTCACC TGGCTGCTC-3', SEQ ID NO:21, corresponding to nucleotides 62-42 of SEQ ID NO:9. The resulting amplification products were analyzed using agarose gel electrophoresis. Five different PCR products were obtained from the five amplification reactions. A DNA fragment of approximately 750 bp which was obtained from the *Ssp*I digested DNA sample was gel extracted. The purified fragment was ligated into the plasmid vector pGEM-T Easy (Promega). The resulting recombinant plasmid was designated pGHP6905 and the nucleotide sequence of the *hpa* insert was determined.

A partial sequence of 594 nucleotides is shown in SEQ ID NO:16. The last nucleotide in SEQ ID NO:13 corresponds to nucleotide 93 in SEQ ID:13. The DNA sequence in SEQ ID NO:16 contains the 5' region of the predicted to contain the promoter region of the *hpa* gene

EXAMPLE 8

Expression of the 592 amino acids HPA polypeptide in a human 293 cell line

The 592 amino acids open reading frame (SEQ ID NOs:13 and 15) was constructed by ligation of the 110 bp corresponding to the 5' end of the SK-hep1 *hpa* cDNA with the placenta cDNA. More specifically the Marathon RACE - PCR amplification product of the placenta *hpa* DNA was digested with *Sac*I and an approximately 1 kb fragment was ligated into a *Sac*I-digested pGHP6905 plasmid. The resulting plasmid was digested with *Ear*I and *Aat*II. The *Ear*I sticky ends were blunted and an approximately 280 bp *Ear*I/blunt-*Aat*II fragment was isolated. This fragment was ligated with pFast*hpa* digested with *Eco*RI which was blunt ended using Klenow fragment and further digested with *Aat*II. The resulting plasmid contained a 1827 bp insert which includes an open reading frame of 1776 bp, 31 bp of 3' UTR and 21 bp of 5' UTR. This plasmid was designated pFastL*hpa*.

A mammalian expression vector was constructed to drive the expression of the 592 amino acids heparanase polypeptide in human cells. The *hpa* cDNA was excised prom pFastL*hpa* with *Bss*HII and *Not*I. The resulting 1850 bp *Bss*HII-*Not*I fragment was ligated to a mammalian expression vector pSI (Promega) digested with *Mlu*I and *Not*I. The

Transient expression of the 592 amino-acids heparanase was examined by western blot analysis and the enzymatic activity was tested using the gel shift assay. Both these procedures are described in length in U.S. Pat. application No. 09/071,739, filed May 1, 1998, which is incorporated by reference as if fully set forth herein. Cells were harvested 3 days following transfection. Harvested cells were re-suspended in lysis buffer containing 150 mM NaCl, 50 mM Tris pH 7.5, 1% Triton X-100, 1 mM PMSF and protease inhibitor cocktail (Boehringer Mannheim). 40 μg protein extract samples were used for separation on a SDS-PAGE. Proteins were transferred onto a PVDF Hybond-P membrane (Amersham). The membrane was incubated with an affinity purified polyclonal anti heparanase antibody, as described in U.S. Pat. application No. 09/071,739. A major band of approximately 50 kDa was observed in the transfected cells as well as a minor band of approximately 65 kDa. A similar pattern was observed in extracts of cells transfected with the pS*hpa* as demonstrated in U.S. Pat. application No. 09/071,739. These two bands probably represent two forms of the recombinant heparanase protein produced by the transfected cells. The 65 kDa protein probably represents a heparanase precursor, while the 50 kDa protein is suggested herein to be the processed or mature form.

pS*hpa*Met- transfected cells was tested by gel shift assay. Cell extracts

transfected and of mock transfected cells were incubated overnight with heparin (6 μg in each reaction) at 37 °C, in the presence of 20 mM phosphate citrate buffer pH 5.4, 1 mM $CaCl_2$, 1 mM DTT and 50 mM NaCl. Reaction mixtures were then separated on a 10 % polyacrylamide gel. The catalytic activity of the recombinant heparanase was clearly demonstrated by a faster migration of the heparin molecules incubated with the transfected cell extract as compared to the control. Faster migration indicates the disappearance of high molecular weight heparin molecules and the generation of low molecular weight degradation products.

EXAMPLE 9

Chromosomal localization of the hpa gene

Chromosomal mapping of the *hpa* gene was performed utilizing a panel of monochromosomal human/CHO and human/mouse somatic cell hybrids, obtained from the UK HGMP Resource Center (Cambridge, England).

40 ng of each of the somatic cell hybrid DNA samples were subjected to PCR amplification using the *hpa* primers: hpu565 5'-AGCTCTGTAGATGTGC TATACAC-3', SEQ ID NO:22, corresponding to nucleotides 564-586 of SEQ ID NO:9 and an antisense

The PCR program was as follows: a hot start of 94 °C – 3 minutes, followed by 7 cycles of 94 °C – 45 seconds, 66 °C – 1 minute, 68 °C – 5 minutes, followed by 30 cycles of 94 °C – 45 seconds, 62 °C – 1 minute, 68 °C – 5 minutes, and a 10 minutes final extension at 72 °C.

The reactions were performed with Expand long PCR (Boehringer Mannheim). The resulting amplification products were analyzed using agarose gel electrophoresis. As demonstrated in Figure 14, a single band of approximately 2.8 Kb was obtained from chromosome 4, as well as from the control human genomic DNA. A 2.8 kb amplification product is expected based on amplification of the genomic *hpa* clone (data not shown). No amplification products were obtained neither in the control DNA samples of hamster and mouse nor in somatic hybrids of other human chromosome.

EXAMPLE 10

Human genomic clone encoding heparanase

Five plaques were isolated following screening of a human genomic library and were designated I.3-1, I.5-1, I.8-1, I.10-1 and I.6-1. The phage DNAs were analyzed by Southern hybridization and by PCR with *hpa* specific and vector specific primers. Southern analysis was performed with three fragments of *hpa* cDNA: a *Pvu*II-*Bam*HI fragment (nucleotides

SEQ ID NO:9) and an *Nde*I-*Xho*I fragment (nucleotides 1103-1721, SEQ ID NO:9).

Following Southern analysis, phages L3, L6, L8 were selected for further analysis. A scheme of the genomic region and the relative position of the three phage clones is depicted in Figure 15. A 2 kb DNA fragment containing the gap between phages L6 and L3 was PCR amplified from human genomic DNA with two gene specific primers GHpuL3 and GHplL6. The PCR product was cloned into the plasmid vector pGEM-T-easy (Promega).

Large scale DNA sequencing of the three Lambda clones and the amplified fragment was performed with Lambda purified DNA by primer walking. A nucleotide sequence of 44,898 bp was analyzed (Figure 16, SEQ ID NO:42). Comparison of the genomic sequence with that of *hpa* cDNA revealed 12 exons separated by 11 introns (Figures 15 an 16). The genomic organization of the *hpa* gene is depicted in Figure 15 (top). The sequence include the coding region from the first ATG to the stop codon which spans 39,113 nucleotides. 2742 nucleotides upstream of the first ATG and 3043 nucleotides downstream of the stop codon. Splice site consensus sequences were identified at exon/intron junctions.

EXAMPLE 11

Alternative splicing

Several minor RT-PCR products were obtained from various cell types, following amplification with *hpa* specific primers. Each one found to contain a deletion of one or two exons. Some of these PCR products contain ORFs, which encode potential shorter proteins.

Table 1 below summarizes the alternative spliced products isolated from various cell lines.

Fragments of similar sizes were obtained following amplification with two cell lines, placenta and platelets.

Cell type	Nucleotides deleted	Exons deleted	ORF
Platelets	1047-1267	8, 9	+
Platelets	1154-1267	9	-
Platelets	289-435, 562-735	2, 4	-
Sk-hep1, platelets, Zr75	562-735	4	+
Sk-hep1 (hepatoma)	561-904	4, 5	-
Zr75 (breast carcinoma)	96-203	1 (partial)	+

EXAMPLE 12

Mouse and rat hpa

EST databases were screened for sequences homologous to the *hpa* gene. Three mouse EST's were identified (accession No. Aa177901, from mouse spleen, Aa067997 from mouse skin, Aa47943 from mouse embryo),

195 bp (SEQ ID NO:12). As shown in Figure 13, the coding region is 80 %

similar to the 3' end of the *hpa* cDNA sequence. These EST's are probably cDNA fragments of the mouse *hpa* homolog that encodes for the mouse heparanase.

Searching for consensus protein domains revealed an amino terminal homology between the heparanase and several precursor proteins such as Procollagen Alpha 1 precursor, Tyrosine-protein kinase-RYK, Fibulin-1, Insulin-like growth factor binding protein and several others. The amino terminus is highly hydrophobic and contains a potential trans membrane domain. The homology to known signal peptide sequences suggests that it could function as a signal peptide for protein localization.

The amino acid sequence of human heparanase was used to search for homologous sequences in the DNA and protein databases. Several human EST's were identified, as well as mouse sequences highly homologous to human heparanase. The following mouse EST's were identified AA177901, AA674378, AA67997, AA047943, AA690179, AI122034, all sharing an identical sequence and correspond to amino acids 336-543 of the human heparanase sequence. The entire mouse heparanase cDNA was cloned, based on the nucleotide sequence of the mouse EST's. PCR primers were designed and a Marathon RACE was performed using a Marathon cDNA library from 15 days mouse embryo (Clontech) and from

amplified from mouse embryo Marathon cDNA library. The first cycle of amplification was performed with primers mhpl773 and Ap1 and the second cycle with primers mhpl736 and AP2. A 1.1 kb fragment was then amplified from BL6 Marathon cDNA library. The first cycle of amplification was performed with the primers mhpl152 and Ap1, and the second with mhpl83 and AP2. The combined sequence was homologous to nucleotides 157 - 1702 of the human *hpa* cDNA, which encode amino acids 33-543. The 5' end of the mouse *hpa* gene was isolated from a mouse genomic DNA library using the Genome Walker kit (Clontech). An 0.9 kb fragment was amplified from a *Dra*I digested Genome walker DNA library. The first cycle of amplification was performed with primers mhpl114 and Ap1 and the second with primers mhpl103 and AP2. The assembled sequence (SEQ ID NOs:43, 45) is 2396 nucleotides long. It contains an open reading frame of 1605 nucleotides, which encode a polypeptide of 535 amino acids (SEQ ID NOs:44, 45), 196 nucleotides of 3' untranslated region (UTR), and anupstream sequence which includes the promoter region and the 5'-UTR of the mouse *hpa* cDNA.. According to two promoter predicting programs TSSW and TSSG, the transcription start site is localized to nucleotide 431 of SEQ ID NOs:43, 45, 163 nucleotides upstream of the first ATG codon. The 431 upstream genomic sequence

average homology of 78 % between the nucleotide sequences and 81 % similarity between the deduced amino acid sequences.

Search for *hpa* homologous sequences, using the Blast 2.0 server revealed two EST's from rat: AI060284 (385 nucleotides, SEQ ID NO:46) which is homologous to the amino terminus (68 % similarity to amino acids 12-136) of human heparanase and AI237828 (541 nucleotides, SEQ ID NO:47) which is homologous to the carboxyl terminus (81 % similarity to amino acids 500-543) of human heparanase, and contains a 3'-UTR. A comparison between the human heparanase and the mouse and rat homologous sequences is demonstrated in Figure 17.

EXAMPLE 13

Prediction of heparanase active site

Homology search of heparanase amino acid sequence against the DNA and the protein databases revealed no significant homologies. The protein secondary structure as predicted by the PHD program consists of alternating alpha helices and beta sheets. The fold recognition server of UCLA predicted alpha/beta barrel structure, with under-threshold confidence.

Five of 15 proteins, which were predicted to have most similar folds,

Lactococcus Lactis, 1amy - alpha-amylase from Barley, 1ecea - endocellulase from Acidothermus Cellulolyticus and 1qbc - hexosaminidase alpha chain, glycosyl hydrolase.

Protein homology search using the bioaccelerator pulled out several proteins, including glycosyl hydrolyses such as beta-fructofuranosidase from *Vicia faba* (broad bean) and from potato, lactase phlorizin hydrolase from human, xylanases from *Clostridium thermocellum* and from *Streptomyces halstedii* and cellulase from *Clostridium thermocellum*. Blocks 9.3 database pulled out the active site of glycosyl hydrolases family five, which includes cellulases from various bacteria and fungi. Similar active site motif is shared by several lysosomal acid hydrolases (63) and other glycosyl hydrolases. The common mechanism shared by these enzymes involves two glutamic acid residues, a proton donor and a nucleophile.

Despite the lack of an overall homology between the heparanase and other glycosyl hydolases, the amino acid couple Asp-Glu (NE), which is characteristic of the proton donor of glycosyl hydrolyses of the GH-A clan, was found at positions 224-225 of the human heparanase protein sequence. As in other clan members, this NE couple is located at the end of a β sheet.

Considering the relative location of the proton donor and the nucleophile in these likely [illegible]

Identification of the active site and the amino acids directly involved in hydrolysis opens the way for expression of the defined catalytic domain. In addition, it will provide the tools for rational design of enzyme activity either by modification of the microenviroment or catalytic site itself.

EXAMPLE 14

Expression of hpa antisense in mammalian cell lines

A mammalian expression vector Hpa2Kepedna3 was constructed in order to express *hpa* antisense in mammalian cells. *hpa* cDNA (1.7 kb *Eco*RI fragment) was cloned into the plasmid pCDNA3 in 3'>5' (antisense) orientation. The construct was used to transfect MBT2-T50 and T24P cell lines. 2 x 10^5 cells in 35 mm plates were transfected using the Fugene protocol (Boehringer Mannheim). 48 hours after transfection cells were trypsinized and seeded in six well plates. 24 hours later G418 was added to initiate selection. The number of colonies per 35 mm plate following 3 weeks:

	Antisense	No insert
T24P	15	60
MBT-T50	1	6

The lower number of colonies obtained after transfection with *hpa* antisense, as compared with the control plasmid suggests that the introduction of *hpa* antisense interfere with cell growth. This experiment demonstrates the use of complementary antisense *hpa* DNA sequence to control heparanase expression in cells. This approach may be used to inhibit expression of heparanase *in vivo*, in, for example, cancer cells and in other pathological processes in which heparanase is involved.

EXAMPLE 15

Zoo blot

Hpa cDNA was used as a probe to detect homologous sequences in human DNA and in DNA of various animals. The autoradiogram of the Southern analysis is presented in Figure 18. Several bands were detected in human DNA, which correlated with the accepted pattern according to the genomic *hpa* sequence. Several intense bands were detected in all mammals, while faint bands were detected in chicken. This correlates with the phylogenetic relation between human and the tested animals. The intense bands indicate that *hpa* is conserved among mammals as well as in more genetically distant organisms. The multiple bands patterns suggest that in all animals, like in human, the *hpa* locus occupy large genomic

based on their homology to the human *hpa* reported herein. This conservation was actually found, between the isolated human *hpa* cDNA and the mouse homologue.

EXAMPLE 16

Characterization of the hpa promoter

The DNA sequence upstream of the *hpa* first ATG was subjected to computational analysis in order to localize the predicted transcription start site and to identify potential transcription factors binding sites. Recognition of human PolII promoter region and start of transcription were predicted using the TSSW and TSSG programs. Both programs identified a promoter region upstream of the coding region. TSSW pointed at nucleotide 2644 and TSSG at 2635 of SEQ ID NO:42. These two predicted transcription start sites are located 4 and 13 nucleotides upstream of the longest *hpa* cDNA isolated by RACE.

A *hpa* promoter-GFP reporter vector was constructed in order to investigate the regulation of *hpa* transcription. Two constructs were made, containing 1.8 kb and 1.1 kb of the *hpa* promoter region. The reporter vector was transfected into T50-mouse bladder carcinoma cells. Cells transfected with both constructs exhibited green fluorescence, which

promoter activity, at various conditions and in different cell types and to characterize the factors involved regulation of *hpa* expression.

Although the invention has been described in conjunction with specific embodiments thereof, it is evident that many alternatives, modifications and variations will be apparent to those skilled in the art. Accordingly, it is intended to embrace all such alternatives, modifications and variations that fall within the spirit and broad scope of the appended claims.

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